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(54) Title: OLEFIN POLYMERIZATION CATALYSTS CONTAINING GROUP 8-10 TRANSITION METALS, BIDENTATE LIGANDS, PROCESSES EMPLOYING SUCH CATALYSTS AND POLYMERS OBTAINED THEREFROM			
(57) Abstract			
<p>Methods for preparing olefin polymers, or oligomers catalysts for preparing olefin polymers and bidentate ligand used for preparing these catalysts are disclosed. The polymers can be prepared by containing the corresponding monomers with a Group 8-10 transition metal catalyst. The polymers are suitable for processing in conventional extrusion processes, and can be formed into high barrier sheets or films, or low molecular weight resins for use in synthetic waxes in wax coatings or as emulsions.</p>			

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Title of the Invention

OLEFIN POLYMERIZATION CATALYSTS CONTAINING GROUP 8-10 TRANSITION METALS, BIDENTATE LIG-
ANTDS, PROCESSES EMPLOYING SUCH CATALYSTS AND POLYMERS OBTAINED THEREFROM

5 **Catalysts And Polymers Obtained Therefrom**

Cross-Reference to Related Applications

This Application Claims the benefit under 35 U.S.C. § 119 (e)
10 of Provisional Application Serial No. 60/039945, filed March 10, 1997;
 Provisional Application Serial No. 60/041542, filed March 25, 1997;
 Provisional Application Serial No. 60/042925, filed April 4, 1997; Provisional
 Application Serial No. 60/043406, filed April 4, 1997; Provisional Application
 Serial No. 60/044691, filed April 18, 1997, and Provisional Application
15 Serial No. 60/059372, filed September 18, 1997.

Field of the Invention

The present invention is directed to Group 8-10 transition metal-
20 containing complexes, their use in olefin polymerizations, and to novel
 olefin polymers produced thereby.

Background of the Invention

25 Olefin polymers are used in a wide variety of products, from
 sheathing for wire and cable to film. Olefin polymers are used, for instance,
 in injection or compression molding applications, in extruded films or
 sheeting, as extrusion coatings on paper, for example photographic paper
 and digital recording paper, and the like. Improvements in catalysts have

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made it possible to better control polymerization processes, and, thus, influence the properties of the bulk material. Increasingly, efforts are being made to tune the physical properties of plastics for lightness, strength, resistance to corrosion, permeability, optical properties, and the like, for 5 particular uses. Chain length, polymer branching and functionality have a significant impact on the physical properties of the polymer. Accordingly, novel catalysts are constantly being sought in attempts to obtain a catalytic process for polymerizing olefins which permits more efficient and better controlled polymerization of olefins.

10 Conventional polyolefins are prepared by a variety of polymerization techniques, including homogeneous liquid phase, gas phase, and slurry polymerization. Certain transition metal catalysts, such as those based on titanium compounds (e.g. $TiCl_3$ or $TiCl_4$) in combination with organoaluminum cocatalysts, are used to make linear and linear low density 15 polyethylenes as well as poly- α -olefins such as polypropylene. These so-called "Ziegler-Natta" catalysts are quite sensitive to oxygen and are ineffective for the copolymerization of nonpolar and polar monomers.

Recent advances in non-Ziegler-Natta olefin polymerization catalysis 20 include the following.

L. K. Johnson et al., WO Patent Application 96/23010, disclose the polymerization of olefins using cationic nickel, palladium, iron, and cobalt complexes containing diimine and bisoxazoline ligands. This document also describes the polymerization of ethylene, acyclic olefins, and/or selected cyclic olefins and optionally selected unsaturated acids or esters 25 such as acrylic acid or alkyl acrylates to provide olefin homopolymers or copolymers.

European Patent Application Serial No. 381,495 describes the polymerization of olefins using palladium and nickel catalysts which contain selected bidentate phosphorous containing ligands.

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L. K. Johnson et al., *J. Am. Chem. Soc.*, 1995, 117, 6414, describe the polymerization of olefins such as ethylene, propylene, and 1-hexene using cationic α -diimine-based nickel and palladium complexes. These catalysts have been described to polymerize ethylene to high molecular weight branched polyethylene. In addition to ethylene, Pd complexes act as catalysts for the polymerization and copolymerization of olefins and methyl acrylate.

G.F. Schmidt et al., *J. Am. Chem. Soc.* 1985, 107, 1443, describe a cobalt(III) cyclopentadienyl catalytic system having the structure [C₅Me₅(L)CoCH₂CH₂- μ -H]⁺, which provides for the "living" polymerization of ethylene.

M. Brookhart et al., *Macromolecules* 1995, 28, 5378, disclose using such "living" catalysts in the synthesis of end-functionalized polyethylene homopolymers.

U. Klabunde, U. S. Patents 4,906,754, 4,716,205, 5,030,606, and 5,175,326, describes the conversion of ethylene to polyethylene using anionic phosphorous, oxygen donors ligated to Ni(II). The polymerization reactions were run between 25 and 100°C with modest yields, producing linear polyethylene having a weight-average molecular weight ranging between 8K and 350 K. In addition, Klabunde describes the preparation of copolymers of ethylene and functional group containing monomers.

M. Peuckert et al., *Organomet.* 1983, 2(5), 594, disclose the oligomerization of ethylene using phosphine, carboxylate donors ligated to Ni(II), which showed modest catalytic activity (0.14 to 1.83 TO/s). The oligomerizations were carried out at 60 to 95° C and 10 to 80 bar ethylene in toluene, to produce linear α -olefins.

R.E. Murray, U.S. Patents Nos. 4,689,437 and 4,716,138, describes the oligomerization of ethylene using phosphine, sulfonate donors ligated to Ni(II). These complexes show catalyst activities approximately 15 times greater than those reported with phosphine, carboxylate analogs.

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W. Keim et al., *Angew. Chem. Int. Ed. Eng.* **1981**, *20*, 116, and V.M. Mohring, et al., *Angew. Chem. Int. Ed. Eng.* **1985**, *24*, 1001, disclose the polymerization of ethylene and the oligomerization of α -olefins with aminobis(imino)phosphorane nickel catalysts; G. Wilke, *Angew. Chem. Int. Ed. Engl.* **1988**, *27*, 185, describes a nickel allyl phosphine complex for the polymerization of ethylene.

5 K.A.O. Starzewski et al., *Angew. Chem. Int. Ed. Engl.* **1987**, *26*, 63, and U. S. Patent 4,691,036, describe a series of bis(ylide) nickel complexes, used to polymerize ethylene to provide high molecular weight

10 linear polyethylene.

WO Patent Application 97/02298 discloses the polymerization of olefins using a variety of neutral N, O, P, or S donor ligands, in combination with a nickel(0) compound and an acid.

15 Brown et al., WO 97/17380, describes the use of Pd α -diimine catalysts for the polymerization of olefins including ethylene in the presence of air and moisture.

1 Fink et al., U. S. Patent No., 4,724,273, have described the polymerization of α -olefins using aminobis(imino)phosphorane nickel catalysts and the compositions of the resulting poly(α -olefins).

20 Additional recent developments are described by Sugimura et al., in JP96-84344, JP96-84343, and WO 9738024, and by Yorisue et al., in JP96-70332.

25 Notwithstanding these advances in non-Ziegler-Natta catalysis, there remains a need for efficient and effective Group 8-10 transition metal catalysts for effecting polymerization of olefins. In addition, there is a need for novel methods of polymerizing olefins employing such effective Group 8-10 transition metal catalysts. In particular, there remains a need for Group 8-10 transition metal olefin polymerization catalysts with both improved temperature stability and functional group compatibility. Further, there

30 remains a need for a method of polymerizing olefins utilizing effective

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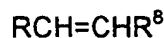
Group 8-10 transition metal catalysts in combination with a Lewis acid so as to obtain a catalyst that is more active and more selective.

Summary of the Invention

5

The present invention is directed to novel Group 8-10 transition metal catalysts, to batch or continuous olefin polymerizations using these catalysts, and to the polymers produced thereby.

10 The process of the present invention comprises contacting one or more olefin monomers of the formula **LI**:



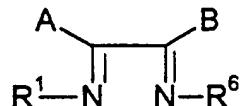
LI

15

wherein R and R⁸ each, independently, represent a hydrogen, a hydrocarbyl or a fluoroalkyl, and may be linked to form a cyclic olefin;

with a Group 8-10 transition metal having coordinated thereto a bidentate ligand having the formula **X**:

20



X

wherein R¹ and R⁶ are each, independently, hydrocarbyl, substituted hydrocarbyl, or silyl, preferably sterically hindered aryl;

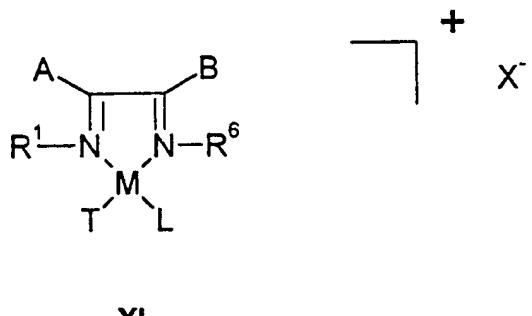
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A and B are each, independently, a heteroatom connected monoradical wherein the connected heteroatom is selected from Group 15 or 16, preferably N, O, S, and wherein A and B may be linked by a bridging group;

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and, optionally, a Bronsted or Lewis acid; and polymerization is carried out to obtain a polymer comprising units of the aforementioned olefin monomer.

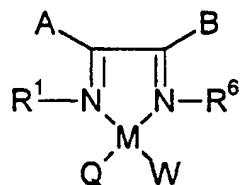
5 In addition, the process of the present invention comprises contacting one or more monomers of the aforementioned formula LI with a catalyst of the present invention having the following formula XI:



10 wherein R¹, R⁶, A, and B are as in formula (X) above;
 T represents a hydrogen or a hydrocarbyl;
 L represents a mono-olefin or a neutral Lewis base wherein the coordinated atom is nitrogen, oxygen, or sulfur;
 M represents a Group 8-10 transition metal, preferably Ni(II), Pd(II),
 15 Co(II) or Fe(II), more preferably Ni(II) or Pd(II);
 and X⁻ is a weakly coordinating anion; and polymerization is carried out to obtain a polymer comprising units of the aforementioned olefin monomer.

Further, the process of the present invention comprises contacting
 20 an olefin monomer of formula LI with a catalyst of the present invention, formed by combining a compound of formula (XII):

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XII

with a compound Y, selected from the group consisting of a neutral Lewis acid capable of abstracting Q⁻ or W to form a weakly coordinating anion, a cationic Lewis acid whose counterion is a weakly coordinating anion, or a Bronsted acid whose conjugate base is a weakly coordinating anion; and

5 wherein R¹, R⁶, A, and B are as in formula X above;

10 Q represents an alkyl, chloride, iodide or bromide;

W represents an alkyl, chloride, iodide or bromide;

M represents a Group 8-10 transition metal, preferably Ni(II), Pd(II), Co(II), or Fe(II); and polymerization is carried out to obtain a polymer comprising units of the aforementioned olefin monomer.

15 The catalysts used in the processes of the present invention readily convert ethylene and α -olefins to high molecular weight polymers, and allow for olefin polymerizations under various conditions, including ambient temperature and pressure, and in solution.

20 The polymers of the present invention include homopolymers of olefins, such as polyethylene, polypropylene, and the like, and to olefin copolymers, including functional-group containing copolymers. As an example, ethylene homopolymers can be prepared with strictly linear to highly branched structures by variation of the catalyst structure, cocatalyst composition, and reaction conditions, including pressure and temperature. The effect these parameters have on polymer structure is described herein.

25 These polymers and copolymers have a wide variety of applications, including use as packaging material and in adhesives.

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Accordingly, it is an object of the present invention to provide novel catalysts capable of polymerizing olefins, including functional group-containing olefins.

It is another object of the invention to set forth catalyst systems for 5 olefin polymerizations.

It is a further object of the invention to describe olefin polymerizations under mild conditions.

It is yet another object of the invention to provide novel catalysts and processes for homogeneous olefin catalysis.

10 It is still a further object of the invention to provide novel polyolefins.

It is yet still another object of the invention to describe methods of varying catalyst structure and polymerization reaction conditions to vary the structure of the resultant polyolefin.

15 These and other objects, features, and advantages of the invention will become apparent as reference is made to the following detailed description and preferred embodiments.

Detailed Description of the Invention

20 In this disclosure certain chemical groups or compounds are described by certain terms and symbols. These terms are defined as follows:

25 Symbols ordinarily used to denote elements in the Periodic Table take their ordinary meaning, unless otherwise specified. Thus, N, O, S, P, and Si stand for nitrogen, oxygen, sulfur, phosphorus, and silicon, respectively.

30 Examples of neutral Lewis acids include, but are not limited to, methylalumoxane (hereinafter MAO) and other aluminum sesquioxides, R^7_3Al , R^7_2AlCl , R^7AlCl_2 (where R^7 is alkyl), organoboron compounds, boron halides, $B(C_6F_5)_3$, BPh_3 , and $B(3,5-(CF_3)C_6H_3)_3$. Examples of ionic

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compounds comprising a cationic Lewis acid include: $R^9_3Sn[BF_4]$, (where R^9 is hydrocarbyl), $MgCl_2$, and H^+X^- , where X^- is a weakly coordinating anion.

Examples of neutral Lewis bases include, but are not limited to, (i) ethers, for example, diethyl ether or tetrahydrofuran, (ii) organic nitriles, for example acetonitrile, (iii) organic sulfides, for example dimethylsulfide, or (iv) monoolefins, for example, ethylene, hexene or cyclopentene.

A "hydrocarbyl" group means a monovalent or divalent, linear, branched or cyclic group which contains only carbon and hydrogen atoms.

Examples of monovalent hydrocarbyls include the following: C_1-C_{20} alkyl; C_1-C_{20} alkyl substituted with one or more groups selected from C_1-C_{20} alkyl, C_3-C_8 cycloalkyl or aryl; C_3-C_8 cycloalkyl; C_3-C_8 cycloalkyl substituted with one or more groups selected from C_1-C_{20} alkyl, C_3-C_8 cycloalkyl or aryl; C_6-C_{14} aryl; and C_6-C_{14} aryl substituted with one or more groups selected from C_1-C_{20} alkyl, C_3-C_8 cycloalkyl or aryl; where the term "aryl" preferably denotes a phenyl, napthyl, or anthracenyl group. Examples of divalent (bridging) hydrocarbyls include: $-CH_2-$, $-CH_2CH_2-$, $-CH_2CH_2CH_2-$, and 1,2-phenylene.

A "silyl" group refers to a SiR_3 group wherein Si is silicon and R is hydrocarbyl or substituted hydrocarbyl or silyl, as in $Si(SiR_3)_3$.

A "heteroatom" refers to an atom other than carbon or hydrogen. Preferred heteroatoms include oxygen, nitrogen, phosphorus, sulfur, selenium, arsenic, chlorine, bromine, silicon and fluorine.

A "substituted hydrocarbyl" refers to a monovalent or divalent hydrocarbyl substituted with one or more heteroatoms. Examples of monovalent substituted hydrocarbyls include: 2,6-dimethyl-4-methoxyphenyl, 2,6-diisopropyl-4-methoxyphenyl, 4-cyano-2,6-dimethylphenyl, 2,6-dimethyl-4-nitrophenyl, 2,6-difluorophenyl, 2,6-dibromophenyl, 2,6-dichlorophenyl, 4-methoxycarbonyl-2,6-dimethylphenyl, 2-tert-butyl-6-chlorophenyl, 2,6-dimethyl-4-phenylsulfonylphenyl, 2,6-

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dimethyl-4-trifluoromethylphenyl, 2,6-dimethyl-4-trimethylammoniumphenyl (associated with a weakly coordinated anion), 2,6-dimethyl-4-hydroxyphenyl, 9-hydroxyanthr-10-yl, 2-chloronaph-1-yl, 4-methoxyphenyl, 4-nitrophenyl, 9-nitroanthr-10-yl, CH_2OCH_3 , cyano, trifluoromethyl, or 5 fluoroalkyl. Examples of divalent (bridging) substituted hydrocarbils include: 4-methoxy-1,2-phenylene, 1-methoxymethyl-1,2-ethanediyl, 1,2-bis(benzyloxymethyl)-1,2-ethanediyl, or 1-(4-methoxyphenyl)-1,2-ethanediyl.

A "sterically hindered aryl" means (i) a phenyl ring with hydrocarbyl, 10 substituted hydrocarbyl, F, Cl, Br or silyl substituents at both the 2- and 6-positions, optionally substituted elsewhere with hydrocarbyl, substituted hydrocarbyl, F, Cl, Br, silyl, hydroxy, methoxy, nitro, cyano, phenylsulfonyl, CO_2Me , CO_2H , $\text{C}(\text{O})\text{CH}_3$, CF_3 , or fluoroalkyl substituents, (ii) a 2-substituted naph-1-yl ring, optionally substituted elsewhere with hydrocarbyl, 15 substituted hydrocarbyl, F, Cl, Br, silyl, hydroxy, methoxy, nitro, cyano, phenylsulfonyl, CO_2Me , CO_2H , $\text{C}(\text{O})\text{CH}_3$, CF_3 , or fluoroalkyl substituents, (iii) an 9-antracenyl ring, optionally substituted elsewhere with hydrocarbyl, substituted hydrocarbyl, F, Cl, Br, silyl, hydroxy, methoxy, nitro, cyano, phenylsulfonyl, CO_2Me , CO_2H , $\text{C}(\text{O})\text{CH}_3$, CF_3 , or fluoroalkyl substituents, or 20 (iv) an aromatic substituted hydrocarbyl with steric properties functionally equivalent (in the context of this invention) to one or more of the following sterically hindered aryls: 2,6-dimethylphenyl, 2,4,6-trimethylphenyl, 2,6-diisopropylphenyl, 2,6-dimethyl-4-nitrophenyl, 2,6-dimethyl-4-phenylsulfonylphenyl, 2-isopropyl-6-methylphenyl, 2,6-bis(trifluoromethyl)phenyl, 2,6-dimethyl-4-methoxyphenyl, 2-methylnaph-1-yl, 9-antracenyl, 2,6-diclorophenyl, 2,6-dibromophenyl, 2-tert-butyl-6-methylphenyl, 2-trimethylsilylnaph-1-yl, 2-chloro-6-methylphenyl, 4-cyano-2,6-dimethylphenyl, 2,6-diisopropyl-4-methoxyphenyl, 2,4,6-tri-tert-butylphenyl, and 2-chloro-6-tert-butylphenyl.

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A "heteroatom connected mono-radical" refers to a mono-radical group in which a heteroatom serves as the point of attachment. Examples include: NH(2,6-dimethylphenyl) and SPh, where Ph is phenyl. Numerous other examples are given herein.

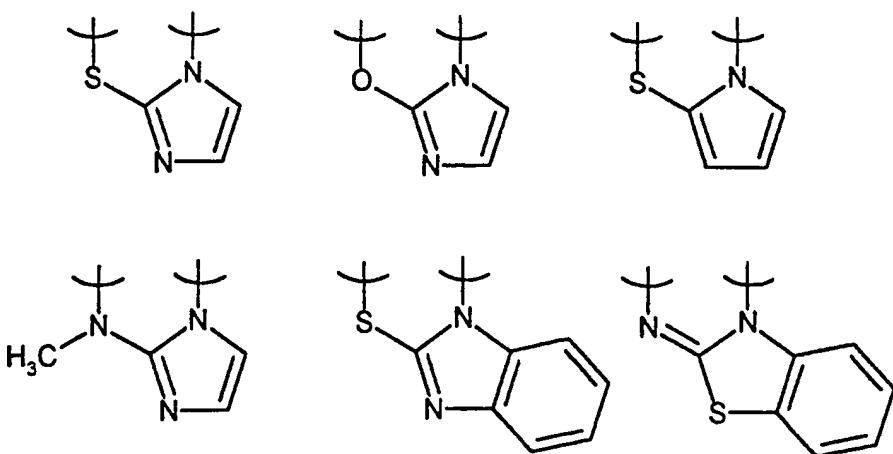
5 A "substituted silicon atom" refers to a $-\text{SiR}^9_2-$ group, wherein R^9 is a hydrocarbyl or substituted hydrocarbyl.

A "substituted phosphorous atom" refers to a $-\text{P}(\text{O})(\text{OR}^9)-$ group, wherein R^9 is a hydrocarbyl or substituted hydrocarbyl.

10 A "substituted sulfur atom" refers to a $-\text{S}(\text{O})-$, $-\text{SO}_2-$, or $-\text{S}(\text{NR}^9)_2-$ group, wherein R^9 is a hydrocarbyl or substituted hydrocarbyl.

A "bridging group" refers to a divalent hydrocarbyl, divalent substituted hydrocarbyl, $-\text{C}(\text{O})-$, $-\text{C}(\text{S})-$, substituted silicon atom, substituted sulfur atom, substituted phosphorous atom, $-\text{CH}_2\text{C}(\text{O})-$, $-\text{C}(\text{O})\text{C}(\text{O})-$, or 3,4,5,6-tetrafluoro-1,2-phenylene.

15 In certain cases, the bridging group, together with groups A and B, may collectively form a divalent heteroatom substituted heterocycle; examples include:



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A "mono-olefin" refers to a hydrocarbon containing one carbon-carbon double bond.

A "suitable metal precursor" refers to a Group 8-10 metal compound, preferably Ni, Co, Pd, and Fe compounds, which may be

5 combined with compound X (preferably, compound III, VI, IX, XVII or XVIII, described below), and optionally a Lewis or Bronsted acid, to form an active olefin polymerization catalyst. Examples include: (1,2-dimethoxyethane)nickel(II) dibromide, bis[(μ -chloro)(1, 2, 3- η ³-2-propenyl)nickel(II)], bis[(μ -chloro)(1, 2, 3- η ³-2-propenyl)palladium(II)], bis[(μ -chloro)(1, 2, 3- η ³-1-trimethylsilyloxy-2-propenyl)nickel(II)], CoBr₂, FeBr₂, 10 bis(acetylacetone)Ni(II), and [tetrakis(acetonitrile)Pd(II)][BF₄].

A "suitable nickel precursor" refers to a suitable metal precursor wherein the metal is nickel.

15 A "suitable nickel(0) precursor" refers to a suitable metal precursor which is a zerovalent nickel compound.

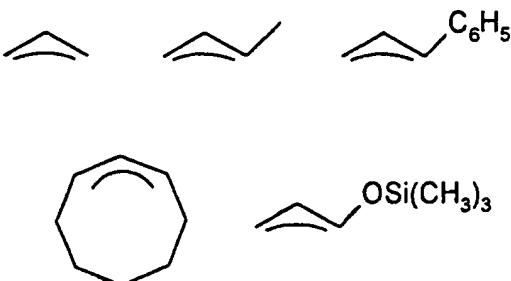
The term "fluoroalkyl" as used herein refers to a C₁-C₂₀ alkyl group substituted by one or more fluorine atoms.

The term "polymer" as used herein is meant a species comprised of monomer units and having a degree of polymerization (DP) of ten or higher.

20 The term " α -olefin" as used herein is a 1-alkene with from 3 to 40 carbon atoms.

A "π-allyl" group refers to a monoanionic group with three sp² carbon atoms bound to a metal center in a η ³-fashion. Any of the three sp² carbon atoms may be substituted with a hydrocarbyl, substituted hydrocarbyl, 25 heteroatom connected hydrocarbyl, heteroatom connected substituted hydrocarbyl, or O-silyl group. Examples of π -allyl groups include:

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where it is understood that the depicted structures are monoanionic fragments.

5 The term π -benzyl group denotes an π -allyl group where two of the sp^2 carbon atoms are part of an aromatic ring. Examples of π -benzyl groups include:



10 where it is understood that the depicted structures are monoanionic fragments.

The term "weakly coordinating anion" is well-known in the art *per se* and generally refers to a large bulky anion capable of delocalization of the negative charge of the anion. Suitable weakly coordinating anions include, 15 but are not limited to, PF_6^- , BF_4^- , SbF_6^- , $(Ph)_4B^-$ wherein Ph = phenyl, tBAr_4 wherein tBAr_4 = tetrakis[3,5-bis(trifluoromethyl)phenyl]borate. The coordinating ability of such anions is known and described in the literature (Strauss, S. et al., *Chem. Rev.* **1993**, *93*, 927).

20 As used herein, the terms "monomer" or "olefin monomer" refer to the olefin or other monomer compound before it has been polymerized; the

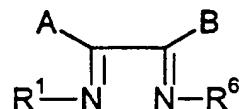
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term "monomer units" refers to the moieties of a polymer that correspond to the monomers after they have been polymerized.

In some cases, a compound Y is required as a cocatalyst. Suitable compounds Y include a neutral Lewis acid capable of abstracting Q⁻ or W to form a weakly coordinating anion, a cationic Lewis acid whose counterion is a weakly coordinating anion, or a Bronsted acid whose conjugate base is a weakly coordinating anion. Preferred compounds Y include: methylalumoxane (hereinafter MAO) and other aluminum sesquioxides, R⁷Al, R⁷AlCl, R⁷AlCl₂ (wherein R⁷ is alkyl), organoboron compounds, boron halides, B(C₆F₅)₃, R⁹Sn[BF₄] (wherein R⁹ hydrocarbyl), MgCl₂, and H⁺X⁻, wherein X⁻ is a weakly coordinating anion.

Examples of "solid support" include inorganic oxide support materials, such as: talcs, silicas, titania, silica/chromia, silica/chromia/titania, silica/alumina, zirconia, aluminum phosphate gels, silanized silica, silica hydrogels, silica xerogels, silica aerogels, silica co-gels.

In one embodiment, the present invention provides a batch or continuous process for the polymerization of olefins, comprising contacting one or more monomers selected from compounds of the formula RCH=CHR⁸ with a catalyst comprising (a) a suitable metal precursor and (b) a ligand of the formula X, and optionally (c) a Bronsted or Lewis acid,



X

wherein R and R⁸ each, independently, represent a hydrogen, a hydrocarbyl, or a fluoroalkyl, and may be linked to form a cyclic olefin;

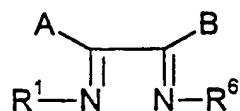
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R^1 and R^6 are each, independently, hydrocarbyl, substituted hydrocarbyl, or silyl, preferably sterically hindered aryl; N represents nitrogen;

5 A and B are each, independently, a heteroatom connected mono-radical wherein the connected heteroatom is selected from Group 15 or 16; in addition, A and B may be linked by a bridging group.

In a further aspect of the invention, there is provided a composition comprising (a) a Group 8-10 transition metal M, (b) one or more Lewis acids, and (c) a binucleating or multinucleating compound of the formula X:

10



X

wherein the Lewis acid or acids are bound to one or more heteroatoms which are π -conjugated to the donor atom or atoms bound to 15 the transition metal M;

15 R^1 and R^6 are each, independently, hydrocarbyl, substituted hydrocarbyl, or silyl, preferably sterically hindered aryl; N represents nitrogen; A and B are each, independently, a heteroatom connected mono-

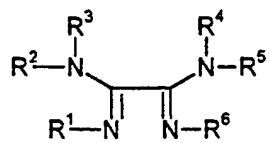
20 radical wherein the connected heteroatom is selected from Group 15 or 16; in addition, A and B may be linked by a bridging group.

Preferred Group 8-10 transition metals are Ni, Pd, Fe, and Co, more preferably Ni and Pd.

Preferred ligands X are those given below:

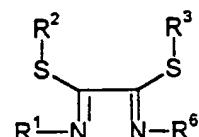
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- 16 -



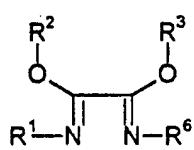
III

or



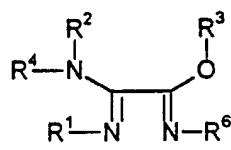
VI

or



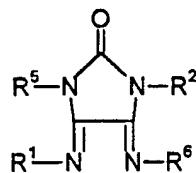
IX

or



XVII

or



XVIII

5

wherein R^1 and R^6 are as defined above;

R^2 , R^3 , R^4 and R^5 each, independently, represent a hydrogen,

hydrocarbyl, substituted hydrocarbyl, or silyl; in addition, any two of R^2 , R^3 ,

R^4 , or R^5 may collectively form a bridging group, provided that when the

10 ligand is of formula VI or IX, the bridging group is not a substituted sulfur atom or a substituted phosphorous atom;

- 17 -

Synthesis of ligands of the formula X in general is based on a modification of methods previously described, and is detailed in the examples given below.

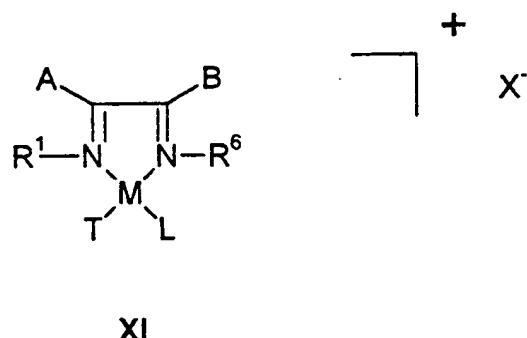
5 The Group 8-10 transition metal complex having coordinated thereto the bidentate ligand of X, above, may also have coordinated to it one or more additional ligands depending, for example, on whether the complex is the active catalyst, a precursor thereto, or the complex is the resting state of the catalyst.

10 In a preferred embodiment of the process of the present invention, one or more olefin monomers of the aforementioned formula LI is contacted with

- (i) a suitable nickel(0) precursor, preferably bis(1,5-cyclooctadiene)nickel(0)compound;
- (ii) a Bronsted acid whose conjugate base is a weakly 15 coordinating anion; and
- (iii) a compound of the aforementioned formula X, preferably a compound of the formula III, VI, IX, XVII, or XVIII, described above; and polymerization is carried out to obtain a polymer comprising units of the aforementioned olefin monomer.

20 In another preferred embodiment of the process of the invention, there is provided a batch or continuous process for the polymerization of olefins, comprising contacting one or more monomers of the formula $RCH=CHR^8$ with a catalyst of formula XI:

- 18 -



wherein R and R⁸ each, independently, represent a hydrogen, a hydrocarbyl, or a fluoroalkyl, and may be linked to form a cyclic olefin;

5 R¹ and R⁶ each, independently, represent hydrocarbyl, substituted hydrocarbyl, or silyl, preferably sterically hindered aryl;

 A and B are each, independently, a heteroatom connected mono-radical wherein the connected heteroatom is selected from Group 15 or 16; in addition, A and B may be linked by a bridging group;

10 T represents a hydrogen or a hydrocarbyl;

 L represents a mono-olefin or a neutral Lewis base wherein the coordinated atom is nitrogen, oxygen, or sulfur;

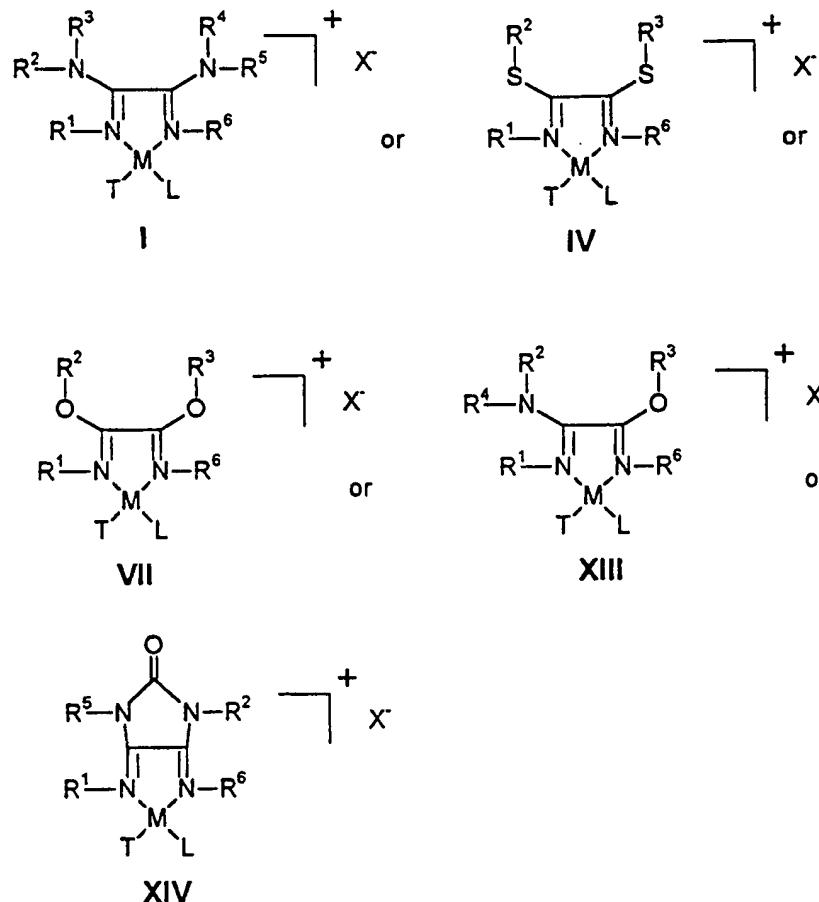
 M represents Ni(II), Pd(II), Co(II), or Fe(II), preferably Ni(II) or Pd(II);

 N represents nitrogen; and

15 X⁻ is a weakly coordinating anion.

In a preferred embodiment, compound XI is selected from compounds having the following formulas:

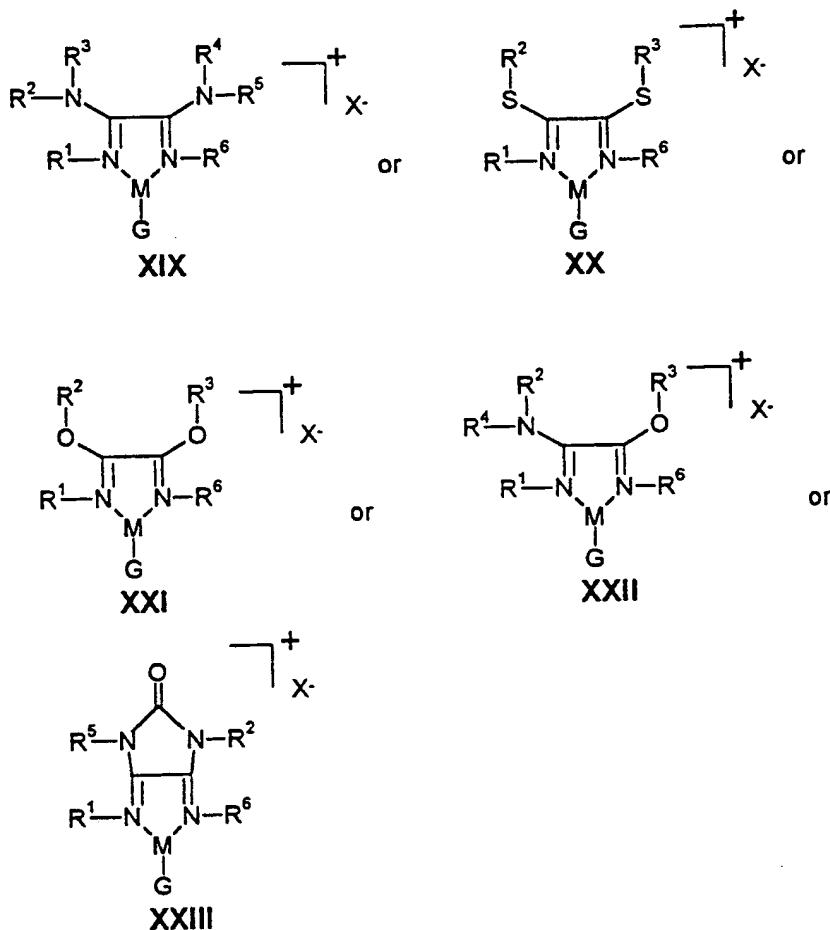
- 19 -



wherein R¹, R², R³, R⁴, R⁵, R⁶, T, L, and M are as defined above.

It is understood that, in the above formulas, when T represents the
 5 growing polymer chain, and L is a mono-olefin, the formulas represent the chain propagating species. T and L may collectively represent a π -allyl or π -benzyl complex, represented by the symbol G in the formula below, which is coordinated to M. Thus, in still another preferred embodiment of the process of the present invention, at least one olefin monomer LI is
 10 contacted with a catalyst selected from the following formulas:

- 20 -



wherein R¹-R⁶, and X⁻ are as defined above;

G is a π -allyl or π -benzyl group;

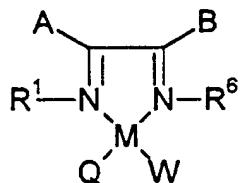
5 M is a Group 8-10 transition metal, preferably Pd(II), Ni(II), Fe(II), or

Co(II), more preferably Ni(II) or Pd(II);

and polymerization is carried out to obtain a polymer comprising units of the aforementioned olefin monomer.

As a further embodiment of the invention, there is provided a batch 10 or continuous process for the polymerization of olefins, comprising contacting one or more monomers of the formula RCH=CHR⁸ with a catalyst formed by combining a compound of formula XII:

- 21 -



XII

with a compound Y,

wherein R and R⁸ each, independently, represent a hydrogen, a hydrocarbyl, or a fluoroalkyl, and may be linked to form a cyclic olefin;

5 R¹ and R⁶ each, independently, represent hydrocarbyl, substituted hydrocarbyl, or silyl, preferably sterically hindered aryl;

A and B are each, independently, a heteroatom connected monoradical wherein the connected heteroatom is selected from Group 15 or 16; in addition, A and B may be linked by a bridging group;

10 Q represents an alkyl, chloride, iodide or bromide;

W represents an alkyl, chloride, iodide or bromide;

N represents nitrogen;

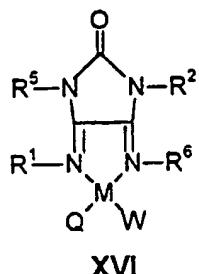
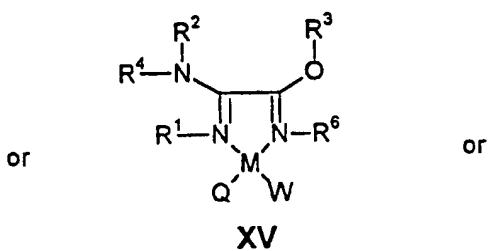
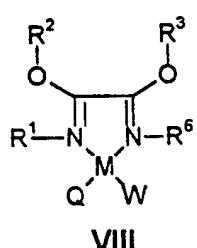
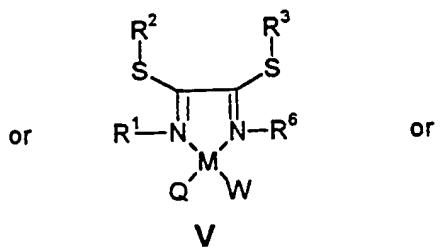
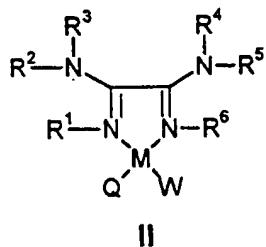
M represents Ni(II), Pd(II), Co(II), or Fe(II), preferably Ni(II) or Pd(II);

and Y is selected from the group consisting of a neutral Lewis acid

15 capable of abstracting Q⁻ or W⁻ to form a weakly coordinating anion, a cationic Lewis acid whose counterion is a weakly coordinating anion, and a Bronsted acid whose conjugate base is a weakly coordinating anion.

In a more preferred embodiment, compounds of formula XII are those selected from the following formulas:

- 22 -



wherein R¹ and R⁶ each, independently, represent hydrocarbyl, substituted hydrocarbyl, or silyl, preferably sterically hindered aryl;

5 R², R³, R⁴ and R⁵ each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl; in addition, any two of R², R³, R⁴, and R⁵ may be linked by a bridging group, provided that when the compound is of formula V or VIII, the bridging group is not a substituted sulfur atom or a substituted phosphorous atom;

10 Q represents an alkyl, chloride, iodide or bromide;

W represents an alkyl, chloride, iodide or bromide;

M represents Ni(II), Pd(II), Co(II) or Fe(II);

- 23 -

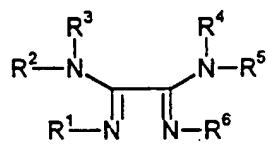
N represents nitrogen; O represents oxygen; S represents sulfur;
and

X⁻ is a weakly coordinating anion.

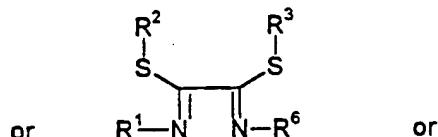
5 As previously mentioned, the catalysts of the present invention may
be used in polymerization processes that incorporate functional group-
containing olefin units within the resultant polymer. Diverse functionalized
olefin polymers and copolymers, and processes for obtaining the same, are
described in copending Provisional Patent Application Serial No. 60-
10 062609, incorporated herein by reference.

The present invention also provides a process for the production of
linear α -olefins, comprising contacting ethylene with (a) a suitable nickel
precursor, (b) a compound of the formula:

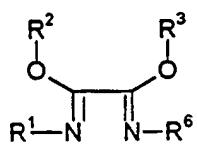
- 24 -



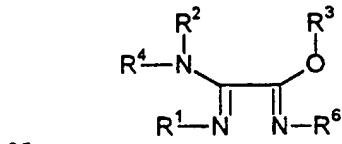
III



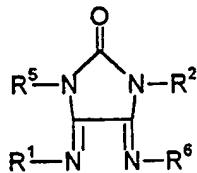
VI



IX



XVII



XVIII

and, optionally (c) a Bronsted or Lewis acid, wherein

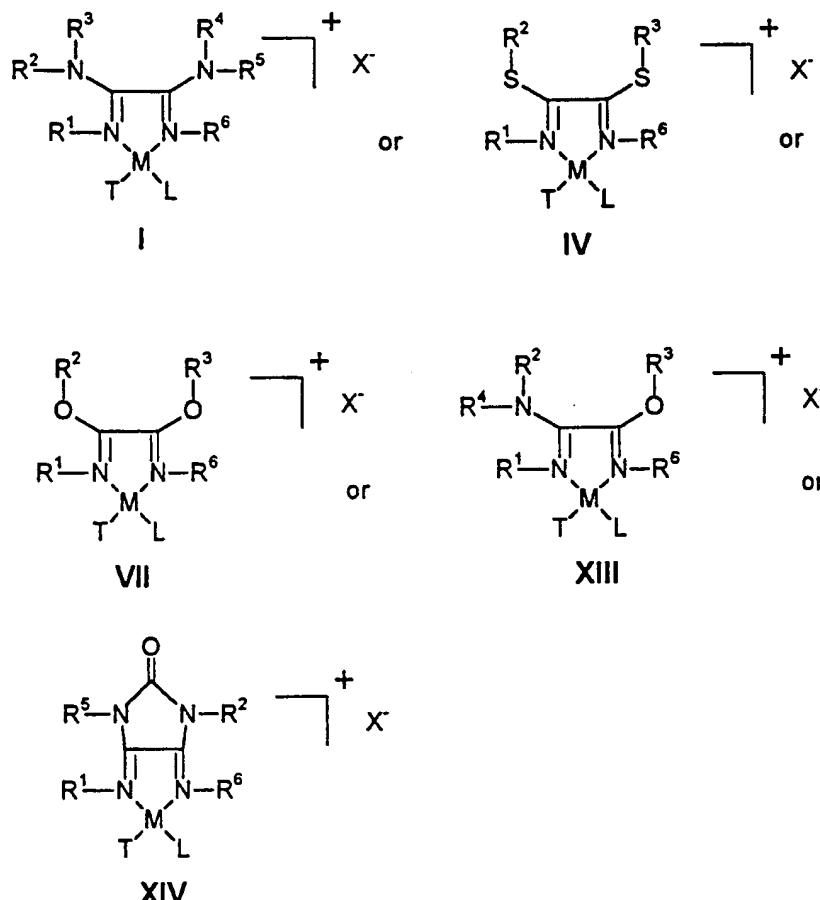
R¹ and R⁶ each, independently, represent hydrocarbyl, substituted hydrocarbyl, or silyl, preferably phenyl or 4-methoxyphenyl;

5 R², R³, R⁴ and R⁵ each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl; in addition, any two of R², R³, R⁴, and R⁵ may collectively form a bridging group;

N represents nitrogen; O represents oxygen; S represents sulfur.

As a further preferred aspect, there is provided a process for the 10 production linear α -olefins, comprising contacting ethylene with a compound of the formula:

- 25 -



wherein R¹ and R⁶ each, independently, represent hydrocarbyl, substituted hydrocarbyl, or silyl, preferably phenyl or 4-methoxyphenyl;

5 R², R³, R⁴ and R⁵ each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl; in addition, any two of R², R³, R⁴, and R⁵ may collectively form a bridging group, provided that when the compound is of formula IV or VII, the bridging group is not a substituted sulfur atom or a substituted phosphorous atom;

10 T represents a hydrogen or hydrocarbyl;

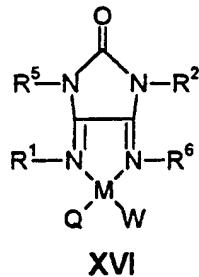
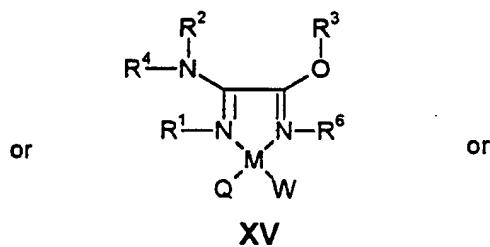
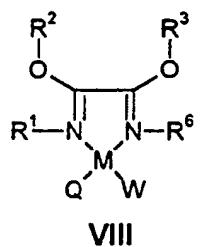
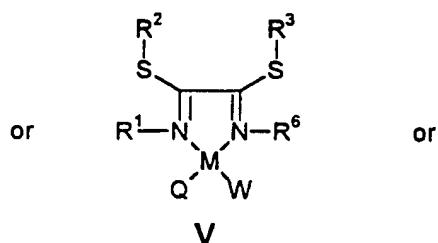
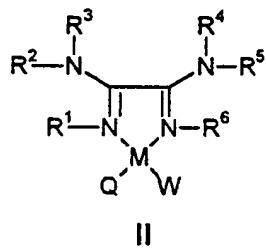
L represents a mono-olefin or a neutral Lewis base wherein the coordinated atom is nitrogen, oxygen, or sulfur;

M represents Ni(II);

- 26 -

N represents nitrogen; O represents oxygen; S represents sulfur;
and
X⁻ is a weakly coordinating anion.

5 As a further preferred aspect, there is provided a process for the production of α -olefins, comprising contacting ethylene with a catalyst formed by combining a complex of the formula:



with a compound Y,

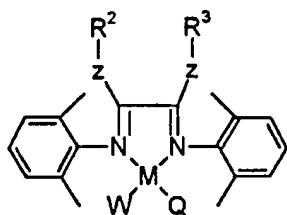
10 wherein R¹ and R⁶ each, independently, represent hydrocarbyl, substituted hydrocarbyl, or silyl, preferably phenyl or 4-methoxyphenyl; R², R³, R⁴ and R⁵ each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl; in addition, any two of R², R³,

- 27 -

R^4 , and R^5 may collectively form bridging group, provided that when the complex is of formula V or VIII, the bridging group is not a substituted sulfur atom or a substituted phosphorous atom;

- 5 Q represents a hydrocarbyl, chloride, iodide or bromide;
- W represents a hydrocarbyl, chloride, iodide or bromide;
- M represents Ni(II);
- N represents nitrogen; O represents oxygen; S represents sulfur;
- 10 and Y is selected from the group consisting of a neutral Lewis acid capable of abstracting Q^- or W^- to form a weakly coordinating anion, a cationic Lewis acid whose counterion is a weakly coordinating anion, or a Bronsted acid whose conjugate base is a weakly coordinating anion.

A further embodiment of the present invention is a process for the copolymerization of ethylene and a comonomer of the formula $CH_2=CH(CH_2)_nCO_2R^1$ which comprises contacting ethylene and a comonomer of the formula $CH_2=CH(CH_2)_nCO_2R^1$ with the complex formed by combining a first compound Y, which is selected from a neutral Lewis acid capable of abstracting Q^- or W^- to form a weakly coordinating anion, a cationic Lewis acid whose counterion is a weakly coordinating anion, or a 20 Bronsted acid whose conjugate base is a weakly coordinating anion; with a second compound of the formula XXIV:



XXIV

wherein R^1 is hydrogen, hydrocarbyl, substituted hydrocarbyl, 25 fluoroalkyl or silyl;

- 28 -

n is an integer greater than 3, preferably greater than 5 and less than 25;

R² and R³ are each independently hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl, or may collectively form a bridging

5 hydrocarbyl, bridging substituted hydrocarbyl, or a substituted silicon atom;

Q is alkyl, chloride, iodide or bromide;

W is alkyl, chloride, iodide or bromide;

N is nitrogen;

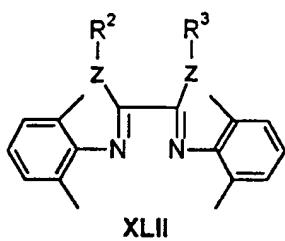
Z is sulfur or oxygen; and

10 M is Ni(II).

As a further aspect of the invention, there is provided a process for the polymerization of ethylene and a comonomer of the formula

CH₂=CH(CH₂)_nCO₂R¹, which comprises contacting ethylene and a

15 comonomer of the formula CH₂=CH(CH₂)_nCO₂R¹ with the complex formed by the reaction product of (a) a compound of formula XLII ; (b) a suitable nickel(0) precursor, and (c) a Bronsted acid whose conjugate base is a weakly coordinating anion,



20

wherein R¹ is hydrogen, hydrocarbyl, substituted hydrocarbyl, fluoroalkyl or silyl;

n is an integer greater than 3, preferably greater than 5 and less than

25;

25 R² and R³ are each independently hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl, or may collectively form a bridging

- 29 -

hydrocarbyl, bridging substituted hydrocarbyl, -C(O)-, or a substituted silicon atom;

N is nitrogen; and

Z is sulfur or oxygen.

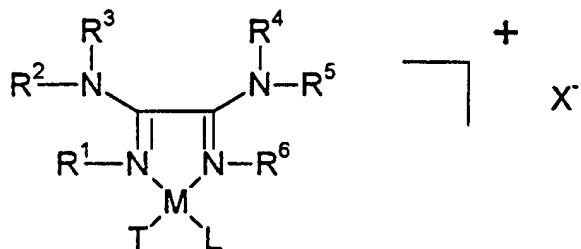
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As a further embodiment of the invention, there is provided a process for the polymerization of olefins comprising contacting one or more monomers of the formula $RCH=CHR^8$ with a binucleating or multinucleating ligand complexed to a Group 8-10 transition metal M and one or more

10 Lewis acids, wherein the Lewis acid or acids are bound to one or more heteroatoms which are π -conjugated to the donor atom or atoms bound to the transition metal M; and R and R^8 each, independently, represent a hydrogen, a hydrocarbyl, or a fluoroalkyl, and may be linked to form a cyclic olefin.

15 The catalysts of the present invention comprise a Group 8-10 transition metal coordinated by the bidentate ligand X; these include, but are not limited to, the preferred compounds I, II, IV, V, VII, VIII, XI-XVI, XIX-XXVIII, and XXX-XLII, set forth in detail herein.

20 A more preferred embodiment of the catalyst of the present invention is a compound of the formula I:



- 30 -

wherein R¹ and R⁶ each, independently, represent hydrocarbyl, substituted hydrocarbyl, or silyl, preferably phenyl, 4-methoxyphenyl, or sterically hindered aryl;

5 R², R³, R⁴ and R⁵ each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl; in addition, any two of R², R³, R⁴, and R⁵ may collectively form a bridging group;

T represents a hydrogen or hydrocarbyl;

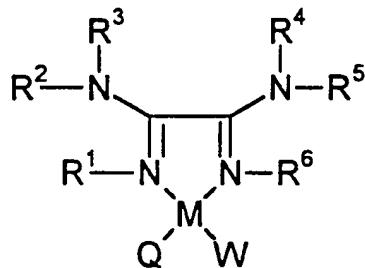
L represents a mono-olefin or a neutral Lewis base wherein the coordinated atom is nitrogen, oxygen, or sulfur;

10 M represents a Group 8-10 transition metal, preferably Ni(II), Pd(II), Co(II), or Fe(II), more preferably Ni(II) or Pd(II);

N represents nitrogen; and

X⁻ is a weakly coordinating anion.

15 Also disclosed is a catalyst comprising a compound of formula II:



II

wherein R¹ and R⁶ each, independently, represent hydrocarbyl, substituted hydrocarbyl, or silyl, preferably phenyl, 4-methoxyphenyl, or sterically hindered aryl;

20 R², R³, R⁴ and R⁵ each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl; in addition, any two of R², R³, R⁴, and R⁵ may collectively form a bridging group;

Q represents a hydrocarbyl, chloride, iodide or bromide;

- 31 -

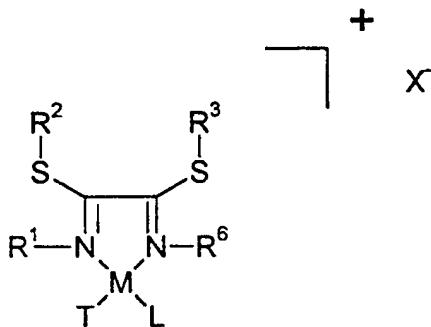
W represents a hydrocarbyl, chloride, iodide or bromide;

N represents nitrogen; and

M represents a Group 8-10 transition metal, preferably Ni(II), Pd(II),

Co(II), or Fe(II), more preferably Ni(II) or Pd(II).

5 Also described is a catalyst comprising a compound of formula IV:



IV

wherein R¹ and R⁶ each, independently, represent hydrocarbyl, substituted hydrocarbyl, or silyl, preferably phenyl, 4-methoxyphenyl, or sterically hindered aryl;

10 R² and R³ each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl, and, in addition, may collectively form a bridging hydrocarbyl, bridging substituted hydrocarbyl, or a substituted silicon atom;

T represents a hydrogen or hydrocarbyl;

15 L represents a mono-olefin or a neutral Lewis base wherein the coordinated atom is nitrogen, oxygen, or sulfur;

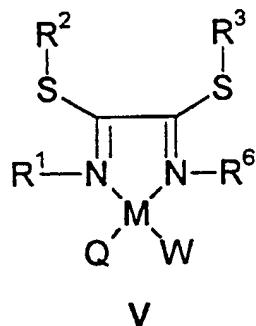
M represents Ni(II), Pd(II), Co(II), or Fe(II), more preferably Ni(II) or Pd(II);

N represents nitrogen; and

20 X is a weakly coordinating anion.

Additionally disclosed is a catalyst comprising a compound of formula V:

- 32 -



wherein R¹ and R⁶ each, independently, represent hydrocarbyl, substituted hydrocarbyl, or silyl, preferably phenyl, or 4-methoxyphenyl or sterically hindered aryl;

5 R² and R³ each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl, and, in addition, may collectively form a bridging hydrocarbyl, bridging substituted hydrocarbyl, or a substituted silicon atom;

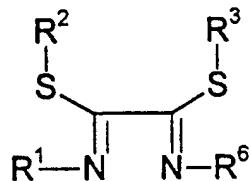
Q represents a hydrocarbyl, chloride, iodide or bromide;

10 W represents a hydrocarbyl, chloride, iodide or bromide;

N represents nitrogen; and

M represents a Group 8-10 transition metal, preferably Ni(II), Pd(II), Co(II), or Fe(II), more preferably Ni(II) or Pd(II);

15 In addition this invention teaches the use of a catalyst comprising a compound of formula VI:



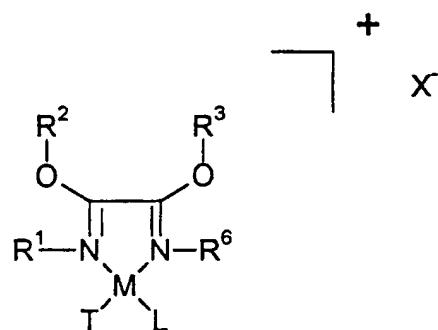
VI

wherein R¹ and R⁶ each, independently, represent hydrocarbyl, substituted hydrocarbyl, or silyl, preferably phenyl, 4-methoxyphenyl, or sterically hindered aryl; and

- 33 -

R^2 and R^3 each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl, and, in addition, may collectively form a bridging hydrocarbyl, bridging substituted hydrocarbyl, or a substituted silicon atom.

5 Described herein is a catalyst comprising a compound of the formula VII:



VII

wherein R¹ and R⁶ each, independently, represent hydrocarbyl, substituted hydrocarbyl, or silyl, preferably phenyl, 4-methoxyphenyl, or 10 sterically hindered aryl;

R^2 and R^3 each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl, and, in addition, may collectively form a bridging hydrocarbyl, bridging substituted hydrocarbyl, or a substituted silicon atom;

15 T represents a hydrogen or hydrocarbyl;

 L represents a mono-olefin or a neutral Lewis base wherein the coordinated atom is nitrogen, oxygen, or sulfur;

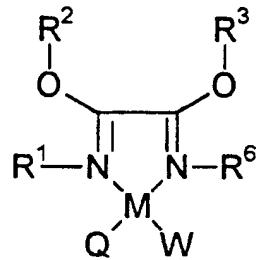
 M represents Ni(II), Pd(II), Co(II), or Fe(II), more preferably Ni(II) or Pd(II);

20 N represents nitrogen; and

 X⁻ is a weakly coordinating anion.

- 34 -

Also described herein is a catalyst comprising a compound of formula VIII:



VIII

wherein R¹ and R⁶ each, independently, represent hydrocarbyl, substituted hydrocarbyl, or silyl, preferably phenyl, 4-methoxyphenyl, or sterically hindered aryl;

R² and R³ each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl, and, in addition, may collectively form a bridging hydrocarbyl, bridging substituted hydrocarbyl, or a substituted silicon atom;

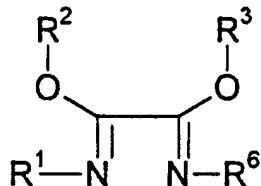
Q represents a hydrocarbyl, chloride, iodide or bromide;

W represents a hydrocarbyl, chloride, iodide or bromide;

N represents nitrogen; and

M represents a Group 8-10 transition metal, preferably Ni(II), Pd(II), Co(II), or Fe(II), more preferably Ni(II) or Pd(II);

Also described herein is a catalyst comprising a compound of formula IX:



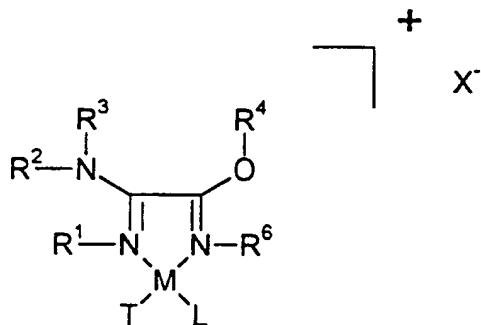
IX

- 35 -

wherein R¹ and R⁶ each, independently, represent hydrocarbyl, substituted hydrocarbyl, or silyl, preferably phenyl, 4-methoxyphenyl, or sterically hindered aryl; and

5 R² and R³ each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl, and, in addition, may collectively form a bridging hydrocarbyl, bridging substituted hydrocarbyl, or a substituted silicon atom.

Also described herein is a catalyst comprising a compound of the formula XIII:



10 XIII

wherein R¹ and R⁶ each, independently, represent hydrocarbyl, substituted hydrocarbyl, or silyl, preferably phenyl, 4-methoxyphenyl, or sterically hindered aryl;

15 R², R³, and R⁴ each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl, and, in addition, may collectively form a bridging group;

T represents a hydrogen or hydrocarbyl;

L represents a mono-olefin or a neutral Lewis base wherein the coordinated atom is nitrogen, oxygen, or sulfur;

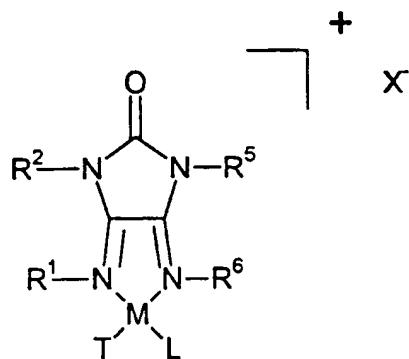
20 M represents Ni(II), Pd(II), Co(II), or Fe(II), more preferably Ni(II) or Pd(II);

N represents nitrogen; and

- 36 -

X^- is a weakly coordinating anion.

Also provided herein is a catalyst comprising a compound of the formula XIV:



XIV

5 wherein R^1 and R^6 each, independently, represent hydrocarbyl, substituted hydrocarbyl, or silyl, preferably phenyl, 4-methoxyphenyl, or sterically hindered aryl;

10 R^2 and R^5 each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl;

10 T represents a hydrogen or hydrocarbyl;

10 L represents a mono-olefin or a neutral Lewis base wherein the coordinated atom is nitrogen, oxygen, or sulfur;

15 M represents a Group 8-10 transition metal, preferably Ni(II), Pd(II), Co(II), or Fe(II), more preferably Ni(II) or Pd(II);

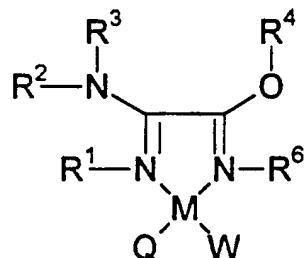
15 N represents nitrogen; and

15 X^- is a weakly coordinating anion.

Also provided herein is a catalyst comprising a compound of formula

XV:

- 37 -



XV

wherein R¹ and R⁶ each, independently, represent hydrocarbyl, substituted hydrocarbyl, or silyl, preferably phenyl, 4-methoxyphenyl, or sterically hindered aryl;

5 R², R³, and R⁴ each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl, and, in addition, may collectively form a bridging group;

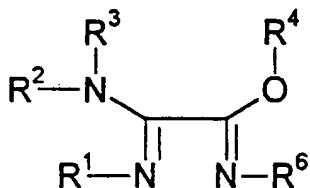
Q represents a hydrocarbyl, chloride, iodide or bromide;

W represents a hydrocarbyl, chloride, iodide or bromide;

10 N represents nitrogen; and

 M represents a Group 8-10 transition metal, preferably Ni(II), Pd(II), Co(II), or Fe(II), preferably Ni(II) or Pd(II).

Also described herein is a catalyst comprising a compound of
15 formula XVII



XVII

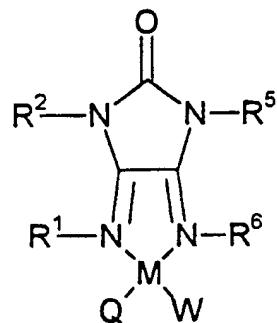
wherein R¹ and R⁶ each, independently, represent hydrocarbyl, substituted hydrocarbyl, or silyl, preferably phenyl, 4-methoxyphenyl, or sterically hindered aryl; and

- 38 -

R^2 , R^3 , and R^4 each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl; in addition, any two of R^2 , R^3 , or R^4 may collectively form a bridging group.

Also provided herein is a catalyst comprising a compound of formula

5 **XVI:**



XVI

wherein R^1 and R^6 each, independently, represent hydrocarbyl, substituted hydrocarbyl, or silyl, preferably phenyl, 4-methoxyphenyl, or sterically hindered aryl;

R^2 and R^5 each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl, preferably sterically hindered aryl;

Q represents a hydrocarbyl, chloride, iodide or bromide;

W represents a hydrocarbyl, chloride, iodide or bromide;

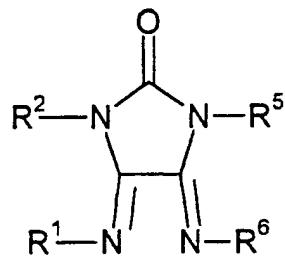
15 N represents nitrogen; and

M represents a Group 8-10 transition metal, preferably Ni(II), Pd(II), Co(II), or Fe(II), more preferably Ni(II) or Pd(II);

Also provided herein is a catalyst comprising a compound of formula

XVIII:

- 39 -

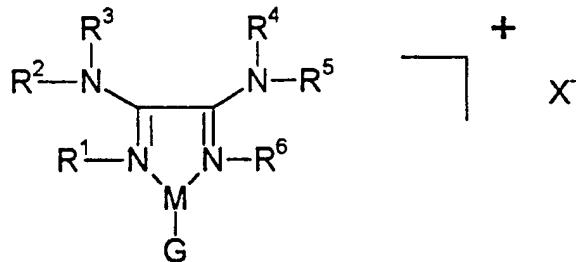


XVIII

wherein R¹ and R⁶ each, independently, represent hydrocarbyl, substituted hydrocarbyl, or silyl, preferably phenyl, 4-methoxyphenyl, or sterically hindered aryl;

5 R² and R⁵ each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl.

Also provided herein is a catalyst comprising a compound of the formula XIX:



XIX

10 wherein R¹ and R⁶ each, independently, represent hydrocarbyl, substituted hydrocarbyl, or silyl, preferably phenyl, 4-methoxyphenyl, or sterically hindered aryl;

15 R², R³, R⁴ and R⁵ each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl; in addition, any two of R², R³, R⁴, and R⁵ may collectively form a bridging group;

G is a π -allyl or π -benzyl group;

M represents Ni(II) or Pd(II);

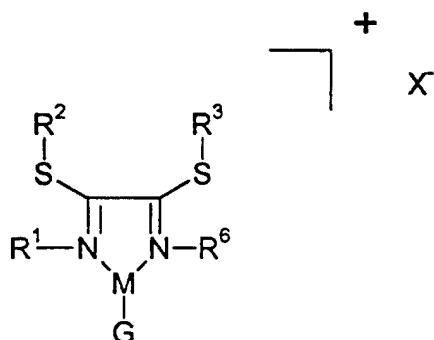
- 40 -

N represents nitrogen; and

X^- is a weakly coordinating anion.

Also provided herein is a catalyst comprising a compound of formula

XX:



XX

5

wherein R¹ and R⁶ each, independently, represent hydrocarbyl, substituted hydrocarbyl, or silyl, preferably phenyl, 4-methoxyphenyl, or sterically hindered aryl;

10 R² and R³ each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl, and, in addition, may collectively form a bridging hydrocarbyl, bridging substituted hydrocarbyl, or a substituted silicon atom;

G is a π -allyl or π -benzyl;

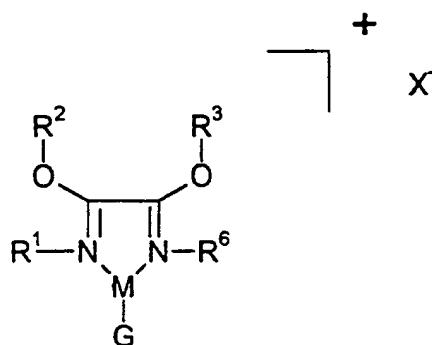
M represents Ni(II) or Pd(II);

15 N represents nitrogen; and

X^- is a weakly coordinating anion.

Also provided herein is a catalyst comprising a compound of the formula **XXI**:

- 41 -



XXI

wherein R¹ and R⁶ each, independently, represent hydrocarbyl, substituted hydrocarbyl, or silyl, preferably phenyl, 4-methoxyphenyl, or sterically hindered aryl;

5 R² and R³ each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl, and, in addition, may collectively form a bridging hydrocarbyl, bridging substituted hydrocarbyl, or a substituted silicon atom;

G is a π -allyl or π -benzyl;

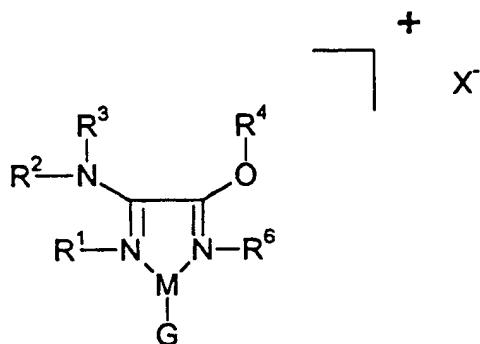
10 M represents Ni(II) or Pd(II);

N represents nitrogen; and

X⁻ is a weakly coordinating anion.

Also provided herein is a catalyst comprising a compound of the formula XXII:

- 42 -



wherein R¹ and R⁶ each, independently, represent hydrocarbyl, substituted hydrocarbyl, or silyl, preferably phenyl, 4-methoxyphenyl, or sterically hindered aryl;

5 R², R³, and R⁴ each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl; in addition, any two of R², R³, and R⁴ may collectively form a bridging group;

G is a π -allyl or π -benzyl;

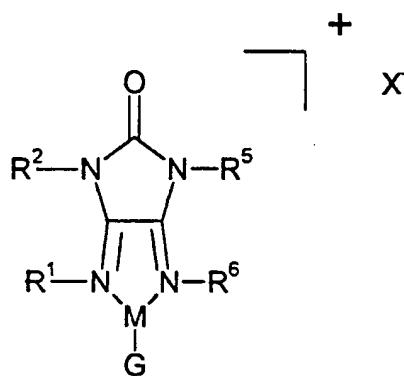
M represents Ni(II) or Pd(II);

10 N represents nitrogen; and

X⁻ is a weakly coordinating anion.

Also provided herein is a catalyst comprising a compound of the formula XXIII:

- 43 -



XXIII

wherein R¹ and R⁶ each, independently, represent hydrocarbyl, substituted hydrocarbyl, or silyl, preferably phenyl, 4-methoxyphenyl, or sterically hindered aryl;

5 R² and R⁵ each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl;

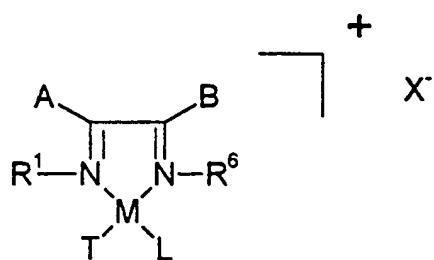
G is a π -allyl or π -benzyl;

M represents Ni(II) or Pd(II);

N represents nitrogen; and

10 X⁻ is a weakly coordinating anion.

Also provided is a compound of formula XXIV,



XXIV

- 44 -

wherein R¹ and R⁶ each, independently, represent hydrocarbyl, substituted hydrocarbyl, or silyl, preferably phenyl, 4-methoxyphenyl, or sterically hindered aryl;

5 A and B are each, independently, a heteroatom connected mono-radical wherein the connected heteroatom is selected from Group 15 or 16; in addition, A and B may be linked by a bridging group;

10 T represents a hydrogen or a hydrocarbyl;

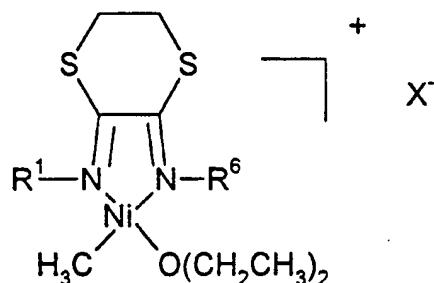
15 L represents a mono-olefin or a neutral Lewis base wherein the coordinated atom is nitrogen, oxygen, or sulfur;

20 M represents Ni(II) or Pd(II);

 N represents nitrogen; and

 X⁻ is a weakly coordinating anion.

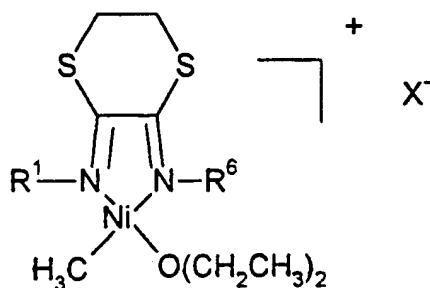
Also provided herein is a catalyst for the polymerization of olefins, 15 comprising a compound of formula XXV,



wherein R¹ and R⁶ are 2,6-dimethylphenyl;
and X⁻ is a weakly coordinating anion.

Also provided herein is a catalyst for the polymerization of olefins, 20 comprising a compound of formula XXVI,

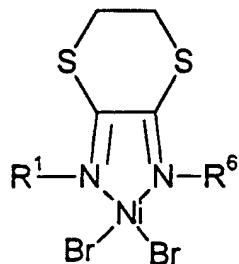
- 45 -



XXVI

wherein R¹ and R⁶ are 2,6-diisopropylphenyl;
and X⁻ is a weakly coordinating anion.

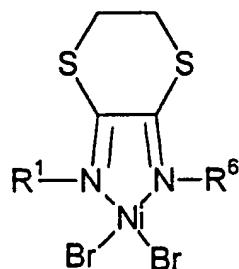
Also provided herein is a catalyst for the polymerization of olefins,
5 comprising a compound of formula XXVII,



XXVII

wherein R¹ and R⁶ are 2,6-dimethylphenyl.

Also provided herein is a catalyst for the polymerization of olefins,
comprising a compound of formula XXVIII,



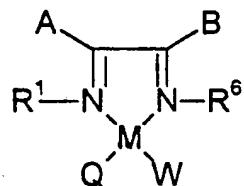
XXVIII

10

wherein R¹ and R⁶ are 2,6-diisopropylphenyl.

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Also provided herein is a catalyst comprising a compound of formula **XXIX**,



XXIX

5

wherein R¹ and R⁶ each, independently, represent hydrocarbyl, substituted hydrocarbyl, or silyl, preferably phenyl, 4-methoxyphenyl, or sterically hindered aryl;

10 A and B are each, independently, a heteroatom connected monoradical wherein the connected heteroatom is selected from Group 15 or 16; in addition, A and B may be linked by a bridging group;

Q represents a hydrocarbyl, chloride, iodide or bromide;

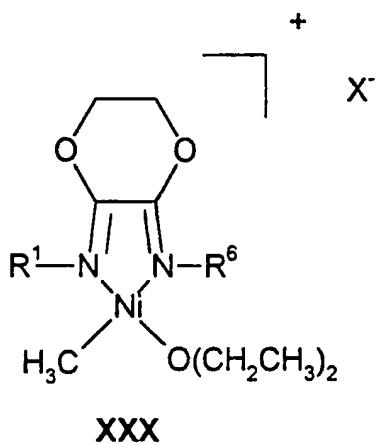
W represents a hydrocarbyl, chloride, iodide or bromide;

M represents Ni(II) or Pd(II); and

15 N represents nitrogen.

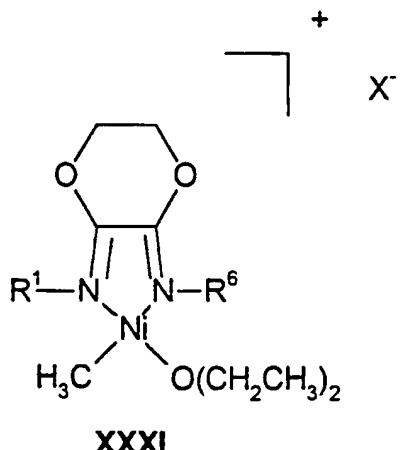
Also provided herein is a catalyst for the polymerization of olefins, comprising a compound of formula **XXX**,

- 47 -



wherein R^1 and R^6 are 2,6-dimethylphenyl;
and X^- is a weakly coordinating anion.

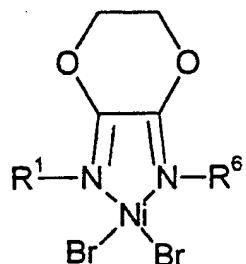
Also provided herein is a catalyst for the polymerization of olefins,
5 comprising a compound of formula XXXI,



wherein R^1 and R^6 are 2,6-diisopropylphenyl;
and X^- is a weakly coordinating anion.

Also provided herein is a catalyst for the polymerization of olefins,
10 comprising a compound of formula XXXII,

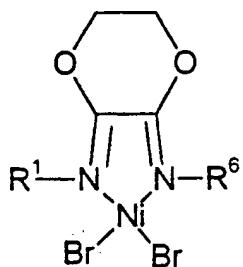
- 48 -



XXXII

wherein R¹ and R⁶ are 2,6-dimethylphenyl.

Also provided herein is a catalyst for the polymerization of olefins, comprising a compound of formula XXXIII,

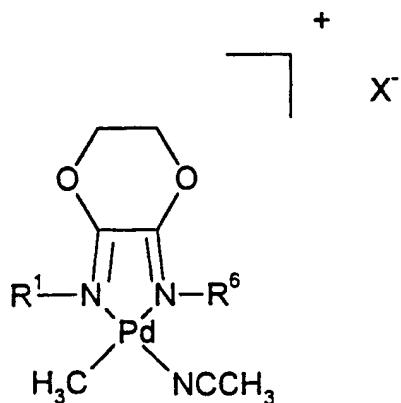


XXXIII

5

wherein R¹ and R⁶ are 2,6-diisopropylphenyl.

Also provided herein is a catalyst for the polymerization of olefins, comprising a compound of formula XXXIV,

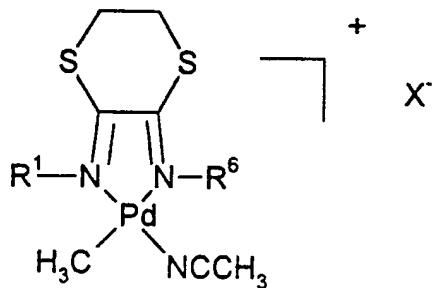


XXXIV

- 49 -

wherein R¹ and R⁶ are 2,6-diisopropylphenyl;
and X⁻ is a weakly coordinating anion.

Also provided herein is a catalyst for the polymerization of olefins,
comprising a compound of formula XXXV,

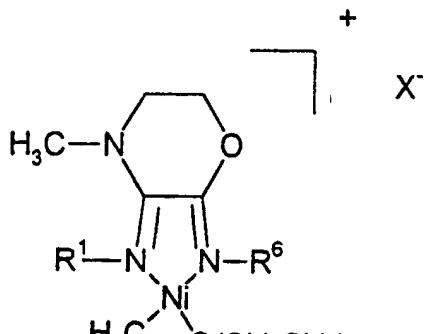


XXXV

5

wherein R¹ and R⁶ are 2,6-diisopropylphenyl;
and X⁻ is a weakly coordinating anion.

Also provided herein is a catalyst for the polymerization of olefins,
comprising a compound of formula XXXVI,



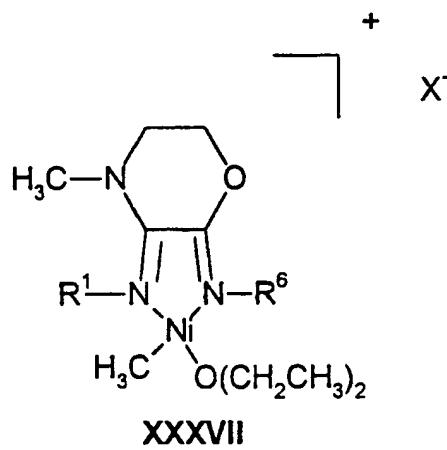
XXXVI

10

wherein R¹ and R⁶ are 2,6-dimethylphenyl;
and X⁻ is a weakly coordinating anion.

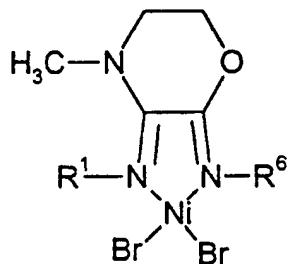
Also provided herein is a catalyst for the polymerization of olefins,
comprising a compound of formula XXXVII,

- 50 -



wherein R¹ and R⁶ are 2,6-diisopropylphenyl;
and X⁻ is a weakly coordinating anion.

Also provided herein is a catalyst for the polymerization of olefins,
5 comprising a compound of formula XXXVIII,

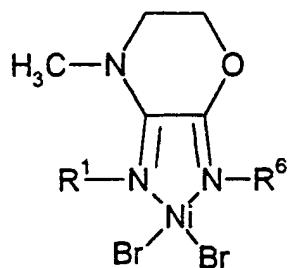


XXXVIII

wherein R¹ and R⁶ are 2,6-dimethylphenyl.

Also provided herein is a catalyst for the polymerization of olefins,
comprising a compound of formula XXXIX,

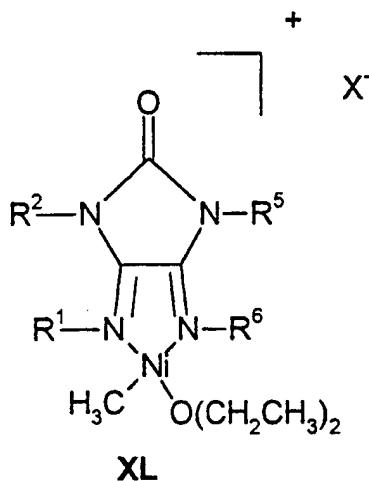
- 51 -



XXXIX

wherein R¹ and R⁶ are 2,6-diisopropylphenyl.

Also provided herein is a catalyst for the polymerization of olefins, comprising a compound of formula XL,

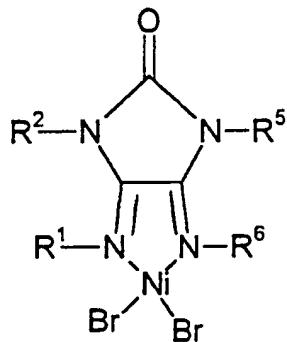


5

wherein R¹, R², R⁵ and R⁶ are 2,6-dimethyl-4-methoxyphenyl; and X⁻ is a weakly coordinating anion.

Also provided herein is a catalyst for the polymerization of olefins, comprising a compound of formula XLI,

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XLI

wherein R¹, R², R⁵ and R⁶ are 2,6-dimethyl-4-methoxyphenyl;
and X⁻ is a weakly coordinating anion.

The invention also provides certain novel polymer compositions.

5 Thus, as a further aspect of the invention, there is provided a polymer composition comprised of monomer units derived from ethylene and from 0.1 to 50 wt % of a comonomer of the formula CH₂=CH(CH₂)_nCO₂R¹, wherein R¹ is hydrogen, hydrocarbyl, substituted hydrocarbyl, fluoroalkyl or silyl, and n is an integer between 3 and 18; in addition to the branches 10 attributable to the incorporation of said comonomer, alkyl branches are present in said polymer composition, wherein between 5 and 50 alkyl branches per 1000 carbon atoms are present and the majority of alkyl branches are methyl branches.

In a preferred embodiment, there is provided a composition 15 comprising an ester containing semicrystalline copolyethylene with 5 to 30 alkyl branches/1000 carbon atoms, wherein the majority of alkyl branches are methyl branches, and an average of from about 1 to 50 ester terminated branches per chain resulting from ester comonomer incorporation.

In a further preferred embodiment, there is provided a composition 20 comprising an olefin containing semicrystalline copolyethylene with 5 to 30 alkyl branches/1000 carbon atoms, wherein the majority of alkyl branches are methyl branches, and an average of from about 1 to 50 olefin

terminated branches per chain resulting from linear diene comonomer incorporation.

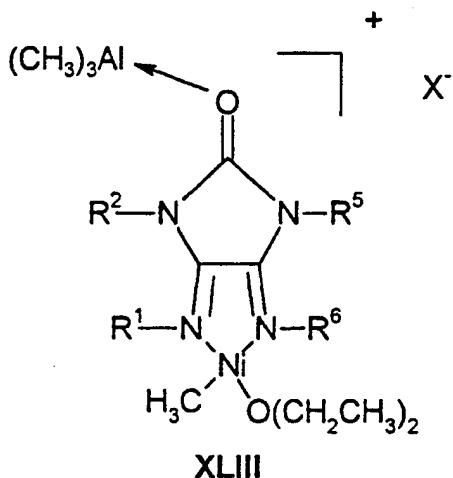
As a further aspect of the invention, there is also provided a polymer composition comprising an ethylene homopolymer with greater than 125 5 alkyl branches per 1000 carbon atoms, useful as an adhesive or tackifying agent.

Preferred olefins useful in the practice of the processes of the present invention include ethylene and α -olefins such as propylene, 1-butene, 1-hexene, 1-octene, ethyl undecenoate and cyclic olefins such as 10 cyclopentene.

When the polymerizations are conducted in the liquid phase, said liquid phase may include solvent or neat monomer. The molar ratio of neutral Lewis acid to transition metal complex can be from 1 to 10000, preferably 10 to 1000. The pressure at which the ethylene polymerizations 15 and copolymerizations take place can be from 1 atmosphere to 1000 atmospheres, preferably 1 to 100 atmospheres.

While not wishing to be bound by theory, the present inventors believe that the Lewis acid may be acting to further activate the catalysts provided herein via coordination to one or more of those heteroatoms which 20 are not directly bound to the transition metal M, but which are π -conjugated to the nitrogens which are bound to the transition metal M. Substituents which contain additional Lewis basic groups, including, but not limited to, methoxy groups, positioned so as to further promote the binding of the Lewis acid at such π -conjugated heteroatoms, are also included in this 25 invention. A nonlimiting example of secondary Lewis acid binding would include the following:

- 54 -



wherein R¹, R², R⁵, and R⁶ are 2,6-dimethylphenyl; and X⁻ is a weakly coordinating anion.

5

The polymerizations may be conducted as solution polymerizations, as non-solvent slurry type polymerizations, as slurry polymerizations using one or more of the olefins or other solvent as the polymerization medium, or in the gas phase. One of ordinary skill in the art, with the present disclosure, would understand that the catalyst could be supported using a suitable catalyst support and methods known in the art. Substantially inert solvents, such as toluene, hydrocarbons, methylene chloride and the like, may be used. Propylene and 1-butene are excellent monomers for use in slurry-type copolymerizations and unused monomer can be flashed off and reused.

10

15

Temperature and olefin pressure have significant effects on copolymer structure, composition, and molecular weight. Suitable polymerization temperatures are preferably from about -100 °C to about 200 °C, more preferably in the 20 °C to 150 °C range.

- 55 -

After the reaction has proceeded for a time sufficient to produce the desired polymers, the polymer can be recovered from the reaction mixture by routine methods of isolation and/or purification.

In general, the polymers of the present invention are useful as 5 components of thermoset materials, as elastomers, as packaging materials, films, compatibilizing agents for polyesters and polyolefins, as a component of tackifying compositions, and as a component of adhesive materials.

High molecular weight resins are readily processed using conventional extrusion, injection molding, compression molding, and 10 vacuum forming techniques well known in the art. Useful articles made from them include films, fibers, bottles and other containers, sheeting, molded objects and the like.

Low molecular weight resins are useful, for example, as synthetic waxes and they may be used in various wax coatings or in emulsion form. 15 They are also particularly useful in blends with ethylene/vinyl acetate or ethylene/methyl acrylate-type copolymers in paper coating or in adhesive applications.

Although not required, typical additives used in olefin or vinyl polymers may be used in the new homopolymers and copolymers of this 20 invention. Typical additives include pigments, colorants, titanium dioxide, carbon black, antioxidants, stabilizers, slip agents, flame retarding agents, and the like. These additives and their use in polymer systems are known *per se* in the art.

The molecular weight data presented in the following examples is 25 determined at 135 °C in 1,2,4-trichlorobenzene using refractive index detection, calibrated using narrow molecular weight distribution poly(styrene) standards.

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EXAMPLES

Other features of the invention will become apparent in the course of the following descriptions of exemplary embodiments which are given for 5 illustration of the invention and are not intended to be limiting thereof.

Example 1

Preparation of *N,N'*-bis(2,6-dimethylphenyl)oxalamide.

2,6-dimethylaniline, triethylamine, and dichloromethane were dried 10 by passage through basic alumina. A 1 L round bottom flask, equipped with a magnetic stir bar and a 125 mL pressure-equalizing dropping funnel capped by a nitrogen inlet adapter, was charged with 53.38 g of 2,6-dimethylaniline, 250 mL of dichloromethane, and 44.76 g of triethylamine. A solution of 25.34 g of oxalyl chloride in 80 mL of dichloromethane was 15 added dropwise under nitrogen with stirring and ice-bath cooling over 1.2 hours to give a thick paste which had to be occasionally swirled by hand to effect mixing. The mixture was allowed to stir at room temperature for 14 hours, then transferred to a separatory funnel, washed 3 times with water, separated and concentrated under reduced pressure (10 mm Hg) to give 63 20 g of crude solid. The crude product was dissolved in a boiling mixture of 1.30 L of toluene and 2.85 L of absolute ethanol, cooled to room temperature (about 23°C) and diluted with 260 mL of water, then allowed to crystallize for 16 hours. The resultant precipitate was isolated by vacuum filtration, washed with methanol (3 x 100 mL) and dried to give 39.1 g (66%) 25 as white crystals. An additional 9.5 g (16.1%) was recovered from the filtrate by further dilution with approximately 500 mL of water. Field desorption mass spectrometry showed a parent ion peak at 296 m/z. ¹H NMR (300 MHz, CDCl₃, chemical shifts in ppm relative to TMS at 0 ppm): 2.29 (12 p, s), 7.15 (6 p, m), 8.86 (2 p, br s).

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Example 2

Preparation of N, N'-bis(2,6-diisopropylphenyl)oxalamide.

2,6-Diisopropylaniline, triethylamine, and dichloromethane were dried by passage through basic alumina. A 1 L round bottom flask, 5 equipped with a magnetic stir bar and a 125 mL pressure-equalizing dropping funnel capped by a nitrogen inlet adapter, was charged with 34.73 g of 2,6-diisopropylaniline (previously distilled), 180 mL of dichloromethane, and 18.30 g of triethylamine. A solution of 10.57 g oxalyl chloride in 43 mL of dichloromethane was added dropwise under nitrogen with stirring and 10 ice-bath cooling over 38 minutes to give a thick paste which had to be occasionally swirled by hand to effect mixing. The mixture was allowed to stir at room temperature (about 23° C) for 60 hours, then diluted with 700 mL of water to precipitate the product, which was isolated by filtration, washed with water and recrystallized from boiling isopropanol (4 L) to afford 15 22.38 g (66%) of white needles. Field desorption mass spectrometry showed a parent ion peak at 408 m/z. ¹H NMR (500 MHz, CD₂Cl₂, chemical shifts in ppm relative to TMS at 0 ppm): 1.22 (24 p, d, 6.8 Hz), 3.08 (4 p, septet, 6.8 Hz), 7.25 (4 p, d, 7.5 Hz), 7.37 (2 p, t, 7.5 Hz), 8.86 (2 p, br s).

20

Example 3

Preparation of N, N'-bis (4-methoxy-2,6-dimethylphenyl)oxalamide.

Triethylamine and dichloromethane were dried by passage through basic alumina. A 50 mL round bottom flask, equipped with a magnetic stir bar and a small pressure-equalized dropping funnel capped by a nitrogen inlet adapter, was charged with 1.5 g of 4-methoxy-2,6-dimethylphenylamine, 8 25 mL of dichloromethane, and 1.38 g of triethylamine. A solution of 0.39 of oxalyl chloride in 2 mL of dichloromethane was added dropwise under a nitrogen atmosphere with stirring and ice-bath cooling over 35 min. The 30 mixture was allowed to stir at room temperature for 14 hours, then

- 58 -

transferred to a separatory funnel, washed with water, separated and concentrated under reduced pressure (10 mm Hg) to give 1.75 g solids. The crude product was dissolved in 150 mL of boiling absolute ethanol and crystallized upon cooling to room temperature (about 23° C.). The resultant precipitate was isolated by vacuum filtration, and dried to give 1.39 g (86%) as white crystals. Field desorption mass spectrometry showed a parent ion peak at 356 m/z.

Example 4

10 Preparation of *N*¹,*N*²-bis(2,6-dimethylphenyl)oxalodiimidoyl dichloride.

A 1 L round bottom flask was charged with 30.0 g of *N,N*'-bis(2,6-dimethylphenyl)oxalamide, 58.8 g of phosphorous pentachloride and 150 mL of dry toluene, and equipped with a magnetic stir bar and a reflux condenser capped by a nitrogen inlet adapter connected to a bubbler. The mixture was heated to reflux over 30 minutes, then maintained at reflux under nitrogen for another 95 minutes to give a yellow solution. Heating was discontinued and the mixture was allowed to cool to room temperature (about 23° C). A short path distillation adapter and receiving flask was attached in place of the condenser and the volatiles were removed under reduced pressure (1 mm Hg), initially at room temperature, then at 100° C, to give 20.1 g (60%) of a granular yellow solid. Field desorption mass spectrometry showed a parent ion peak at 332 m/z. ¹H NMR (300 MHz, C₆D₆, chemical shifts in ppm relative to TMS at 0 ppm): 2.04 (12 p, s), 6.91 (6 p, s).

25

Example 5

Preparation of *N*¹,*N*²-bis(2,6-diisopropylphenyl)oxalodiimidoyl dichloride.

A 500 mL round bottom flask equipped with a magnetic stir bar and a reflux condenser capped by a nitrogen inlet adapter connected to a bubbler was charged with 2.50 g of *N,N*'-bis(2,6-

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diisopropylphenyl)oxalamide, 3.58 g of phosphorous pentachloride and 36 mL of dry toluene. The mixture was heated to reflux over 30 minutes, then maintained at reflux under nitrogen for another 210 minutes to afford a clear yellow solution. Heating was discontinued and the mixture was allowed to 5 cool to room temperature (about 23° C). A short path distillation adapter and receiving flask were attached in place of the condenser and the volatiles were removed under reduced pressure (1 mm Hg), initially at room temperature, then at 100° C, to give a yellow oil, which slowly crystallized upon complete cooling. The product was purified by column 10 chromatography (SiO₂, Merck Grade 9385 230-400 mesh, 60 Å; 3 v% ethyl acetate in hexane) to afford 1.49 g (55%) yellow crystals. Field desorption mass spectrometry showed a parent ion peak at 444 m/z.

Example 6

15 Preparation of *N¹,N²*-bis(4-methoxy-2,6-dimethylphenyl)oxalodiimidoyl dichloride.

A 50 mL round bottom flask was charged with 1.37 g of *N,N*'-bis(4-methoxy-2,6-dimethylphenyl)oxalamide 1.88 g of phosphorous pentachloride and 15 mL of dry toluene, and equipped with a magnetic stir bar and a reflux 20 condenser capped by a nitrogen inlet adapter connected to a bubbler. The mixture was heated, with stirring, at about 100° C until the evolution of HCl ceased. Then, another 0.22 g PCls was added and the mixture was heated another 30 min at 80° C. After cooling to room temperature the mixture was transferred to a separatory funnel, and some crystallization occurred upon 25 transfer. Complete transfer and re-dissolution of the product was accomplished by the addition of dichloromethane. The organic layer was washed with saturated aqueous sodium bicarbonate, then concentrated *in vacuo* to afford 1.44 g crystalline yellow solid. Field desorption mass spectrometry showed a parent ion peak at 392 m/z.

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Example 7

Preparation of 2,3-bis(2,6-dimethylphenylimino)-1,4-dithiane.

A 50 mL round bottom flask equipped with a magnetic stir bar and a reflux condenser capped by a nitrogen inlet was charged with 504 mg of N^1,N^2 -bis(2,6-dimethylphenyl)oxalodiimidoyl dichloride, 136 mg of sodium hydride (60% mineral oil dispersion), 4.0 mL of dry tetrahydrofuran and 0.140 mL of 1,2-ethanedithiol. The mixture was heated at reflux for 2 hours, after which another 66 mg of sodium hydride dispersion was added and the mixture was refluxed for an additional hour. After cooling, the mixture was diluted with water and diethyl ether, and the ether layer was separated, washed again with water, and dried with magnesium sulfate to afford a yellow-orange oil. Column chromatography (SiO₂, Merck Grade 9385 230-400 mesh, 60 Å; 15 v% of ethyl acetate in hexane) afforded 296 mg of a yellow oil which was crystallized by addition of hexane and collected by vacuum filtration to give 219 mg of yellow granular crystals. Field desorption mass spectrometry showed a parent ion peak at 354 m/z. ¹H NMR (300 MHz, CDCl₃, chemical shifts in ppm relative to TMS at 0 ppm): 2.17 (12p, s), 3.27 (4p, br s), 6.4-7.12 (6 p, m).

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Example 8

Preparation of 2,3-bis(2,6-diisopropylphenylimino)-[1,4]dithiane.

A 250 mL round bottom flask equipped with a magnetic stir bar and a reflux condenser capped by an argon inlet was sequentially charged with 0.69 g of a 60% dispersion of sodium hydride in mineral oil, 3.34 g of N^1,N^2 -bis(2,6-diisopropylphenyl)oxalodiimidoyl dichloride, 20 mL of dry tetrahydrofuran, and 0.70 mL of 1,2-ethanedithiol. The mixture was heated under argon at reflux for 3 hours, after which another 0.25 g of sodium hydride dispersion was added and the mixture was refluxed for an additional 2.5 hours. After cooling, the mixture was diluted with water and diethyl ether, and the ether layer was separated, washed with water, and

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dried with magnesium sulfate to afford a yellow-orange oil. Column chromatography (SiO₂, Merck Grade 9385 230-400 mesh, 60 Å; 10 v% ethyl acetate in hexane) afforded 3.14 g of a yellow-orange glass. Field desorption mass spectrometry showed a parent ion peak at 466 m/z. ¹H NMR (500 MHz, CD₂Cl₂, chemical shifts in ppm relative to TMS at 0 ppm): 1.10-1.22 (12 p, m), 1.22-1.40 (12 p, m), 2.78-3.05 (4 p, m), 3.30 (4 p, br s), 7.05-7.25 (6 p, m).

Example 9

10 Preparation of 2,3-bis(phenylimino)-[1,4]dithiane.

A 250 mL round bottom flask equipped with a magnetic stir bar and a reflux condenser capped by an argon inlet was sequentially charged with 0.69 g of a 60% dispersion of sodium hydride in mineral oil, a freshly prepared solution of 2.08 g *N*¹,*N*²- diphenyloxalodiimidoyl dichloride in 20 mL of dry tetrahydrofuran, and 0.70 mL of 1,2-ethanedithiol. The mixture was heated under argon at reflux for 2 hours, after which another 0.30 g of sodium hydride dispersion was added and the mixture refluxed for an additional 3 hours. After cooling, the mixture was diluted with water and diethyl ether, and the ether layer was separated, washed with water, and dried with magnesium sulfate to afford a yellow-orange gummy solid. Column chromatography (SiO₂, Merck Grade 9385 230-400 mesh, 60 Å; 10 v% ethyl acetate in hexane) afforded 296 mg of a yellow oil which was crystallized by addition of hexane to give 0.161 g of yellow-orange granular crystals. Field desorption mass spectrometry showed a parent ion peak at 298 m/z. ¹H NMR (300 MHz, CDCl₃, chemical shifts in ppm relative to TMS at 0 ppm): 3.27 (4p, br s), 7.02 (4p, apparent d, 8.1 Hz), 7.19 (2p, apparent t, 7.2 Hz), 7.40 (4p, apparent t, 7.8 Hz).

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Example 10

Preparation of 2,3-bis(2,6-dimethylphenylimino)-2,3-dihydrobenzo[1,4]dithiine.

A 100 mL round bottom flask equipped with a magnetic stir bar and a reflux condenser capped by an argon inlet was sequentially charged with 0.294 g of a 60% dispersion of sodium hydride in mineral oil, 4 mL of dry tetrahydrofuran, and 0.253 g of 1,2-benzenedithiol. After the bubbling had subsided, 0.600 g of N^1,N^2 -bis(2,6-dimethylphenyl)oxalodiimidoyl dichloride was added. The mixture was stirred at 25° C. for 45 minutes, then heated to reflux over 15 minutes and held at reflux for 1 hour. After cooling, the mixture was diluted with water and diethyl ether, and the ether layer was separated, washed with water, and dried with magnesium sulfate, and concentrated *in vacuo* to give a yellow-orange oil. Column chromatography (SiO₂, Merck Grade 9385 230-400 mesh, 60 Å; 2 v% ethyl acetate in hexane) afforded 0.412 g of a yellow-orange glass. Field desorption mass spectrometry showed a parent ion peak at 402 m/z. ¹H NMR (300 MHz, CDCl₃, chemical shifts in ppm relative to TMS at 0 ppm): 2.16 (12 p, s), 7.01-7.24 (10 p, m).

Example 11

Preparation of 2,3- bis(4-methoxy-2,6-dimethylphenylimino)-[1,4]dithiane.

A 50 mL round bottom flask equipped with a magnetic stir bar and a reflux condenser capped by a nitrogen inlet was charged with 420 mg of N^1,N^2 -bis(4-methoxy-2,6-dimethylphenyl)oxalodiimidoyl dichloride. To a 50 mL pear flask was added 171 mg of sodium hydride (60% mineral oil dispersion), 1.75 mL of dry tetrahydrofuran, and, cautiously, 0.11 mL ethane dithiol. The resulting mixture was syringed into the N^1,N^2 -bis(4-methoxy-2,6-dimethylphenyl)oxalodiimidoyl dichloride solution using 5 mL dry THF to complete the transfer. The reaction flask was heated at reflux for 3 hours, after which another 45 mg of sodium hydride dispersion and

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another 20 μ L ethane dithiol was added and the mixture was refluxed for an additional hour. After cooling, the mixture was diluted with water and diethyl ether, and the ether layer was separated, washed again with water, dried with magnesium sulfate, and concentrated to afford a yellow-orange solid. Column chromatography (SiO₂, Merck Grade 9385 230-400 mesh, 60 \AA ; 15 v% ethyl acetate in hexane) afforded 147 mg of a yellow powder. Field desorption mass spectrometry showed a parent ion peak at 414 m/z.

Example 12

10 Preparation of 2,3-bis(2,6-dimethylphenylimino)-[1,4]dioxane.

A 50 mL round bottom flask equipped with a magnetic stir bar and a reflux condenser capped by a nitrogen inlet was charged with 504 mg of N¹,N²-bis(2,6-dimethylphenyl)oxalodiimidoyl dichloride, 66 mg of sodium hydride (60% mineral oil dispersion), 5.0 mL of dry tetrahydrofuran, 0.230 mL of triethylamine (dried by passage through alumina) and 0.093 mL of dry ethylene glycol. The mixture was heated at reflux for 105 minutes, after which another 66 mg of sodium hydride dispersion was added and the mixture was refluxed for an additional hour. After cooling, the mixture was diluted with water and diethyl ether, and the ether layer was separated, washed again with water, dried with magnesium sulfate, and concentrated to afford a yellow oil. Crystallization from heptane gave rosettes of off-white crystals (225 mg, 1st crop). Field desorption mass spectrometry showed a parent ion peak at 322 m/z. ¹H NMR (300 MHz, CDCl₃, chemical shifts in ppm relative to TMS at 0 ppm): 2.20 (12 p, s), 4.35 (4 p, s), 6.94 (2 p, m), 7.05 (4 p, m).

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Example 13

Preparation of 5-methoxymethyl-2,3-bis(2,6-dimethylphenylimino)-[1,4]dioxane.

A 50 mL round bottom flask equipped with a magnetic stir bar and a reflux condenser capped by an argon inlet was charged with 504 mg of 5 N^1,N^2 -bis(2,6-dimethylphenyl)oxalodiimidoyl dichloride, 155 mg of sodium hydride (60% mineral oil dispersion), 3.5 mL of dry tetrahydrofuran and 188 mg of 3-methoxy-1,2-propanediol. The mixture was heated to reflux over 10 min and held at reflux for 2 hours. After cooling, the mixture was diluted with diethyl ether, and washed with water (2 x 100 mL), and dried with 10 magnesium sulfate and concentrated *in vacuo* to afford 329 mg of a yellow oil. Column chromatography (SiO₂, Merck Grade 9385 230-400 mesh, 60 Å; 20 v% of ethyl acetate in hexane) afforded 216 mg of a glassy yellow solid. Field desorption mass spectrometry showed a parent ion peak at 366 15 m/z. ¹H NMR (300 MHz, CDCl₃, chemical shifts in ppm relative to TMS at 0 ppm): 2.18 (12 p, s), 3.31 (3 p, s), 3.45-3.65 (2 p, m), 4.20-4.40 (2 p, m), 4.40-4.55 (1 p, m), 6.80-7.15 (6 p, m).

Example 14

20 Preparation of 2,3-bis(benzyloxymethyl)-5,6-bis(2,6-dimethylphenylimino)-[1,4]dioxane.

A 100 mL round bottom flask equipped with a magnetic stir bar and a reflux condenser capped by an argon inlet was charged with 265 mg of 25 sodium hydride (60% mineral oil dispersion), 7.5 mL of dry tetrahydrofuran, 1.0 g of 3-methoxy-1,2-propanediol and 1.0 g of N^1,N^2 -bis(2,6-dimethylphenyl)oxalodiimidoyl dichloride. The yellow mixture was heated to reflux over 15 min and became very viscous. More tetrahydrofuran (5 mL) was added and the mixture was stirred with a glass rod, then heated for another 30 min. Next, an additional 220 mg of sodium hydride (60% 30 mineral oil dispersion) and an additional 10 mL tetrahydrofuran were added.

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Most of the yellow color was discharged with the second addition of sodium hydride, rendering the very viscous reaction mixture light brown. Heating was continued for about 15 min more, and after cooling, the mixture was diluted with diethyl ether and washed with water to remove the sodium chloride. Column chromatography (SiO₂, Merck Grade 9385 230-400 mesh, 60 Å; 2 v% of ethyl acetate in hexane) afforded a beige gummy solid. Field desorption mass spectrometry showed a parent ion peak at 562 m/z. ¹H NMR (300 MHz, CDCl₃, chemical shifts in ppm relative to TMS at 0 ppm): 2.17 (12 p, s), 3.63 (4 p, br s), 4.38 (2 p, d, 12.6 Hz), 4.47 (2 p, d, 12.6 Hz), 10 4.56 (2 p, br s), 6.85-7.4 (16 p, m).

Example 15

Preparation of 2,3-bis(2,6-diisopropylphenylimino)-[1,4]dioxane.

A 50 mL round bottom flask equipped with a magnetic stir bar and a reflux condenser capped by a nitrogen inlet was charged with 1.0 g of N¹,N²-bis(2,6-diisopropylphenyl)oxalodiimidoyl dichloride, 268 mg of sodium hydride (60% mineral oil dispersion), 4.0 mL of dry tetrahydrofuran, and 212 mg of dry ethylene glycol. Under an argon atmosphere, the mixture was heated to 65° C held at that temperature for 90 minutes. The mixture was then quickly heated to reflux and held at reflux for 30 minutes more. After cooling, the mixture was diluted with 50 mL diethyl ether, and washed with water, dried with magnesium sulfate, and concentrated *in vacuo* to afford a light straw-yellow oil (903 mg). Column chromatography (SiO₂, Merck Grade 9385 230-400 mesh, 60 Å; 8 v% ethyl acetate in hexane) afforded 257 mg of a pale green glass. Field desorption mass spectrometry showed a parent ion peak at 434 m/z. ¹H NMR (300 MHz, CDCl₃, chemical shifts in ppm relative to TMS at 0 ppm): 1.24 (24 p, d, 6.6 Hz), 3.00 (4p, septet, 6.6 Hz), 4.31 (4 p, br s), 7.04-7.20 (6 p, m).

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Example 16

Preparation of 2,3-bis(2,6-dimethylphenylimino)-4-methylmorpholine.

A 50 mL round bottom flask equipped with a magnetic stir bar and a reflux condenser capped by a nitrogen inlet adapter was charged with 503 mg of *N*¹,*N*²-bis(2,6-dimethylphenyl)oxalodiimidoyl dichloride, 346 mg triethylamine, 4 mL dry, deoxygenated toluene, and 0.135 mL 2-(methylamino)ethanol. The mixture was heated to reflux under nitrogen over 30 minutes and maintained at reflux for another 4.25 hours. After cooling, the mixture was diluted with 45 mL diethyl ether and washed three times with water (110 mL total). The ether extract was dried with magnesium sulfate, filtered and concentrated under reduced pressure (10 mm Hg) to give a light-colored solid (512 mg). Recrystallization from heptane/dichloromethane gave 138 mg pale off-white crystals (first crop). Field desorption mass spectrometry showed a parent ion peak at 335 m/z. ¹H NMR (300 MHz, CDCl₃, chemical shifts in ppm relative to TMS at 0 ppm): 1.73 (6 p, s), 2.10 (6 p, s), 3.27 (3 p, s), 3.55-3.65 (2 p, m), 4.16-4.26 (2 p, m), 6.65-6.95 (6 p, m).

Example 17

Preparation of 2,3-bis(2,6-diisopropylphenylimino)-4-methylmorpholine.

A 50 mL round bottom flask equipped with a magnetic stir bar and a reflux condenser capped by a nitrogen inlet adapter was charged with 725 mg of *N*¹,*N*²-bis(2,6-diisopropylphenyl)oxalodiimidoyl dichloride, 143 mg of sodium hydride (60% mineral oil dispersion), 4 mL dry tetrahydrofuran, and 0.144 mL 2-(methylamino)ethanol. The mixture was stirred at room temperature for 4 hours, and allowed to stand at room temperature for another 5 days. The mixture was diluted with diethyl ether and washed with water. The ether extract was concentrated under reduced pressure (10 mm Hg) to give a yellow oil which partially crystallized over several hours). Column chromatography (SiO₂, Merck Grade 9385 230-400 mesh, 60 Å; 12

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v% of ethyl acetate in hexane) afforded 156 mg light yellow crystals. Field desorption mass spectrometry showed a parent ion peak at 447 m/z. ¹H NMR (300 MHz, CDCl₃, chemical shifts in ppm relative to TMS at 0 ppm): 0.86 (6 p, d, 7.2 Hz), 1.04 (6 p, d, 7.2 Hz), 1.15 (6 p, d, 7.2 Hz), 1.18 (6 p, d, 7.2 Hz), 2.27 (2 p, apparent septet, 7.2 Hz), 2.97 (2 p, apparent septet, 7.2 Hz), 3.28 (3 p, br s), 3.55-3.65 (2 p, m), 4.14-4.22 (2 p, m), 6.80-7.02 (6 p, m).

Example 18

10 Preparation of 1,3-bis-(2,6-dimethyl-phenyl)-4,5-bis-(2,6-dimethyl-phenylimino)-imidazolidin-2-one.

In a 250 mL round bottom flask, 1.0 g of *N¹,N²,N³,N⁴-tetrakis(2,6-dimethylphenyl)oxalamidine was dissolved in 35 mL dry, deoxygenated dichloromethane while stirring under an argon atmosphere. 0.67 mL dry triethylamine was added, followed by 240 mg triphosgene. With the addition of the triphosgene, the color shifted from pale yellow to chrome yellow. The mixture was allowed to stir for 16 h at room temperature, after which time an additional 460 mg of triphosgene was added. After about 15 min, 10 mg dimethylamino pyridine and an additional 240 mg triphosgene were added. After about 15 min more, another 220 mg triphosgene and about 0.5 g dimethylamino pyridine were added. The mixture was washed with saturated aqueous sodium bicarbonate, then with water, and then concentrated *in vacuo* to afford a yellow powder. Column chromatography (SiO₂, Merck Grade 9385 230-400 mesh, 60 Å; 4 v% ethyl acetate in hexane) afforded 763 mg of a chrome yellow powder. Field desorption mass spectrometry showed a parent ion peak at 528 m/z. ¹H NMR (300 MHz, CDCl₃, chemical shifts in ppm relative to TMS at 0 ppm): 2.01 (12 p, s), 2.32 (12 p, s), 6.4-7.3 (12 p, m).*

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Example 19

Preparation of 1,3-bis(4-methoxy-2,6-dimethylphenyl)-4,5-bis-(4-methoxy-2,6-dimethylphenylimino)imidazolidin-2-one.

A 100 mL round bottom was equipped with a magnetic stirrer and charged
5 with 0.75 mL dry triethylamine, 6 mL dry and deoxygenated
dichloromethane, and 0.335 g N^1,N^2,N^3,N^4 -tetrakis(4-methoxy-2,6-
dimethylphenyl)oxalamidine. With stirring, 178 mg triphosgene was added,
and the flask was quickly capped with a septum. A precipitate formed and
the color shifted from yellow to orange. After 2.5 days another 78 mg
10 triphosgene was added, and the reaction left to stir another 2 hours. 150
mg more triphosgene was added, and the reaction left to stir for another 16
hrs. The reaction mixture was diluted with 50 mL diethyl ether and washed
with water (2 x 50 mL). The aqueous washings were back-extracted with
dichloromethane. The organic layers were combined and dried over
15 magnesium sulfate, and concentrated *in vacuo* to afford an orange oil.
Upon addition of diethyl ether to the oil, small orange crystals formed. The
compound was isolated on a vacuum filter and washed with diethyl ether to
afford 216 mg yellow-orange microcrystalline powder. Field desorption
mass spectrometry showed a parent ion peak at 648 m/z. 1H NMR (500
20 MHz, CD_2Cl_2 , chemical shifts in ppm relative to TMS at 0 ppm): 1.98 (12 p,
broad hump), 2.25 (12 p, broad hump), 3.63 (6 p, broad hump), 3.75 (6 p,
broad hump), 6.32 (4 p, broad hump), 6.59 (4 p, broad hump).

Example 20

Preparation of N^1,N^2,N^3,N^4 -tetrakis(2,6-dimethylphenyl)oxalamidine.

A 1 L round bottom flask equipped with a magnetic stir bar and a
reflux condenser capped by a nitrogen inlet was charged with 5.6 g of
 N^1,N^2 -bis(2,6-dimethylphenyl)oxalodiimidoyl dichloride, 43 mL of dry
toluene and 32.7 g of 2,6-dimethylaniline (dried by passage through
30 alumina). The mixture was heated to reflux under nitrogen over 30 minutes,

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then maintained at reflux another 3 hours. After cooling, the mixture was diluted with 206 g of absolute ethanol and 45 g of water to produce copious amounts of precipitate. Isolation by vacuum filtration, with ethanol (600 mL) and heptane (600 mL) washes, and subsequent drying, gave 6.1 g (72%)
5 as pale yellow crystals. Field desorption mass spectrometry showed a parent ion peak at 502 m/z. ^1H NMR (300 MHz, CDCl_3 , chemical shifts in ppm relative to TMS at 0 ppm): 2.16 (24 p, s), 6.75 (12 p, s), 8.6 (2 p, br s).

Example 21

10 Preparation of N^1,N^2,N^3,N^4 -tetrakis(4-methoxy-2,6-dimethylphenyl)oxalamidine.

A 500 mL round bottom flask equipped with a magnetic stir bar and a reflux condenser capped by a nitrogen inlet was charged with 1.0 g of N^1,N^2 -bis(4-methoxy-2,6-dimethylphenyl)oxalodiimidoyl dichloride, 24 mL of dry toluene
15 and 854 mg of 4-methoxy-2,6-dimethylphenylamine, and 0.90 mL triethylamine (dried by passage through alumina). The mixture was heated to reflux under nitrogen over 30 minutes, then maintained at reflux another 14 hours. After cooling, the mixture was diluted with dichloromethane and washed with water. The impure compound was adsorbed onto SiO_2 , and
20 column chromatography (SiO_2 , Merck Grade 9385 230-400 mesh, 60 Å; 12.5 v% ethyl acetate in hexane) afforded 340 mg of a yellow powder. Field desorption mass spectrometry showed a parent ion peak at 622 m/z.

Example 22

25 Preparation of 1,4-dimethyl-2,3-bis(2,6-dimethylphenylimino)piperazine.

A 25 mL round bottom flask equipped with a magnetic stir bar and a reflux condenser capped by a nitrogen inlet adapter was charged with 0.50 g of N^1,N^2 -bis(2,6-dimethylphenyl)oxalodiimidoyl dichloride, 1.1 mL of N,N' -dimethylethylenediamine and 4.0 mL of dry toluene. The mixture was
30 heated to reflux under nitrogen over 15 minutes and maintained at reflux for

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another 30 minutes. After cooling, the mixture was diluted with diethyl ether and washed three times with water. The ether extract was dried with magnesium sulfate, filtered and concentrated under reduced pressure (10 mm Hg) to give a yellow solid (0.50 g). Recrystallization from heptane gave 5 pale yellow crystals. Field desorption mass spectrometry showed a parent ion peak at 348 m/z. ^1H NMR (300 MHz, CDCl_3 , chemical shifts in ppm relative to TMS at 0 ppm): 1.83 (br s, 12 p), 2-3.4 (two very broad humps, 4 p), 3.42 (br s, 6 p), 6.66 (t, 2p), 6.84 (d, 4p).

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Example 23

Preparation of the nickel dibromide complex of 2,3-bis(2,6-dimethylphenylimino)-[1,4]dithiane.

A 50 mL Schlenk flask equipped with a magnetic stir bar and capped with a septum was charged with 100 mg of 2,3-bis(2,6-dimethylphenylimino)-[1,4]dithiane and 79 mg of (1,2-dimethoxyethane)nickel(II) dibromide under an inert atmosphere. Dry, deoxygenated dichloromethane (5 mL) was added and the mixture was stirred under an argon atmosphere, turning red-brown within about 5 minutes and slowly producing a red-brown crystalline precipitate. After 1 15 hour, another 5 mL of dichloromethane was added. The mixture was stirred another 21 hours at 21° C, then diluted with 10 mL of dry, deoxygenated hexane and stirred another 8 hours. The supernatant was removed via a filter paper-tipped cannula, and the residue dried *in vacuo* at 1 mm Hg to afford 116 mg of red-brown crystals.

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Example 24

Preparation of the nickel dibromide complex of 2,3-bis(2,6-diisopropylphenylimino)-[1,4]dithiane.

A Schlenk flask equipped with a magnetic stir bar was charged 30 with 79 mg of 2,3-bis(2,6-diisopropylphenylimino)-[1,4]dithiane (0.17 mmol)

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and 49 mg of (1,2-dimethoxyethane)nickel(II) dibromide (0.16 mmol) under an argon atmosphere. Dry, deoxygenated dichloromethane (15 mL) was added and the mixture was stirred under an argon atmosphere, turning red-brown within about 10 minutes. After 2 hours, the CH_2Cl_2 was removed *in vacuo*. The resulting red-brown solid was washed with 2x10 mL of hexane and the solid was dried *in vacuo* for several hours affording 76 mg of a brown solid.

Example 25

10 Preparation of the nickel dibromide complex of 2,3-bis(phenylimino)-[1,4]dithiane.

A 50 mL Schlenk flask equipped with a magnetic stir bar and capped with a septum was charged with 151 mg of 2,3-bis(phenylimino)-[1,4]dithiane and 123 mg of (1,2-dimethoxyethane)nickel(II) dibromide under an inert atmosphere. Dry, deoxygenated dichloromethane (10 mL) was added and the mixture was stirred under an argon atmosphere, turning dark brown within about 5 minutes and slowly producing a red-brown crystalline precipitate. After 80 minutes, the mixture was concentrated to apparent dryness under a stream of argon, then further dried *in vacuo* for 1 hour at 50 mTorr to afford a red-brown powder.

Example 26

Preparation of the nickel dibromide complex of 2,3-bis(2,6-dimethylphenylimino)-2,3-dihydrobenzo[1,4]dithiine.

25 A 50 mL Schlenk flask equipped with a magnetic stir bar and capped with a septum was charged with 110 mg of 2,3-bis-(2,6-dimethylphenylimino)-2,3-dihydrobenzo[1,4]dithiine and 71 mg of (1,2-dimethoxyethane)nickel(II) dibromide under an inert atmosphere. Dry, deoxygenated dichloromethane (8 mL) was added and the mixture was stirred under an argon atmosphere, quickly turning red-brown. The mixture

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was stirred for 1 hour, concentrated to dryness under a stream of argon, then further dried *in vacuo* for 1 hour at 50 mTorr to yield a red-brown crystalline powder.

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Example 27

Preparation of the nickel dibromide complex of 2,3- bis(4-methoxy-2,6-dimethylphenylimino)-[1,4]dithiane.

A 50 mL Schlenk flask equipped with a magnetic stir bar and capped with a septum was charged with 147 mg of 2,3- bis(4-methoxy-2,6-dimethylphenylimino)-[1,4]dithiane and 93 mg of (1,2-dimethoxyethane)nickel(II) dibromide under an inert atmosphere. Dry, deoxygenated dichloromethane (10 mL) was added and the mixture was stirred under an argon atmosphere, turning dark brown almost immediately, and producing a brown precipitate. After 2 hours, 10 mL dry and deoxygenated hexane was added to complete the precipitation. The supernatant was removed via filter paper-tipped cannula, the residue dried *in vacuo* (0.5 mm Hg) for 14 h to obtain the product as a brown, microcrystalline solid.

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Example 28

Preparation of the nickel dibromide complex of 2,3-bis(2,6-dimethylphenylimino)-[1,4]dioxane.

A 50 mL Schlenk flask equipped with a magnetic stir bar and capped with a septum was charged with 100 mg of 2,3-bis(2,6-dimethylphenylimino)-[1,4]dioxane and 87 mg of (1,2-dimethoxyethane)nickel(II) dibromide under an inert atmosphere. Dry, deoxygenated dichloromethane (5 mL) was added and the mixture was stirred under an argon atmosphere, slowly producing a brown crystalline precipitate. After 1 hour, another 5 mL of dichloromethane was added. The mixture was stirred another 21 hours at 21° C, then diluted with 10 mL of

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dry, deoxygenated hexane and stirred another 8 hours. The supernatant was removed via a filter paper-tipped cannula, and the residue dried *in vacuo* at 1 mm Hg to afford 117 mg of brown crystals.

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Example 29

Preparation of the nickel dibromide complex of 2,3-bis(benzylloxymethyl)-5,6-bis(2,6-dimethylphenylimino)-[1,4]dioxane.

A 50 mL Schlenk flask equipped with a magnetic stir bar and capped with a septum was charged with 172 mg of 2,3-bis(benzylloxymethyl)-5,6-bis(2,6-dimethylphenylimino)-[1,4]dioxane and 85 mg of (1,2-dimethoxyethane)nickel(II) dibromide under an inert atmosphere. Dry, deoxygenated dichloromethane (12 mL) was added and the mixture was stirred under an argon atmosphere, almost immediately turning red-brown. After 1.75 hours, the mixture was concentrated dryness under a stream of argon for 16 h, then further dried *in vacuo* to afford 182 mg of a red-brown crystalline powder.

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Example 30

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Preparation of the nickel dibromide complex of 2,3-bis(2,6-dimethylphenylimino)-4-methylmorpholine.

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A 25 mL Schlenk flask equipped with a magnetic stir bar and capped with a septum was charged with 100 mg of 2,3-bis(2,6-dimethylphenylimino)-4-methylmorpholine and 84 mg of (1,2-dimethoxyethane)nickel(II) dibromide under an inert atmosphere. Dry, deoxygenated dichloromethane (5 mL) was added and the mixture was stirred under an argon atmosphere for 1 hour, after which another 5 mL dichloromethane was added. After 16 hours, the mixture was diluted with 10 mL hexane, and the supernatant was removed via a filter paper tipped cannula, and the residue was dried *in vacuo* to obtain 139 mg green crystals.

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Example 31

Reaction of 1,3-bis-(2,6-dimethyl-phenyl)-4,5-bis-(2,6-dimethyl-phenylimino)-imidazolidin-2-one, (1,2-dimethoxyethane)nickel(II) dibromide, 5 and silver tetrafluoroborate.

In an argon filled glove box, a flame-dried Schlenk flask equipped with a magnetic stir bar was charged with 159.6 mg of 1,3-bis-(2,6-dimethyl-phenyl)-4,5-bis-(2,6-dimethyl-phenylimino)-imidazolidin-2-one and 92.7 mg of (1,2-dimethoxyethane)nickel(II) dibromide and 59.4 mg silver 10 tetrafluoroborate. The flask was wrapped in aluminum foil, and on the Schlenk line, under an argon atmosphere, 10 mL dry tetrahydrofuran was added. A white precipitate immediately separated. The mixture was stirred for 25 min, then the supernatant was transferred via filter paper-tipped cannula to a dry septum-capped vial. The supernatant was concentrated to 15 dryness under a stream of dry argon for 16 h to afford 256 mg of a yellow crystalline powder.

Example 32

Preparation of the nickel dibromide complex of 1,3-bis(4-methoxy-2,6-dimethylphenyl)-4,5-bis(4-methoxy-2,6-dimethylphenylimino)imidazolidin-2-one.

A 50 mL Schlenk flask equipped with a magnetic stir bar and capped with a septum was charged with 101 mg of 1,3-bis(4-methoxy-2,6-dimethylphenyl)-4,5-bis(4-methoxy-2,6-dimethylphenylimino)imidazolidin-2-one and 40 mg of (1,2-dimethoxyethane)nickel(II) dibromide under an inert atmosphere. Dry, deoxygenated dichloromethane (10 mL) was added and the mixture was stirred under an nitrogen atmosphere, slowly turning dark red-brown over 3 hours. After 2 more hours, the supernatant was removed 25 to a flame-dried Schlenk flask via a filter paper-tipped cannula, diluted with 30 10 mL dry, deoxygenated hexane, and concentrated to dryness under a

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stream of nitrogen to give a mixture of a tan microcrystalline powder and large, well-defined dark brown crystals. The latter were separated and used without further purification.

5

Example 33

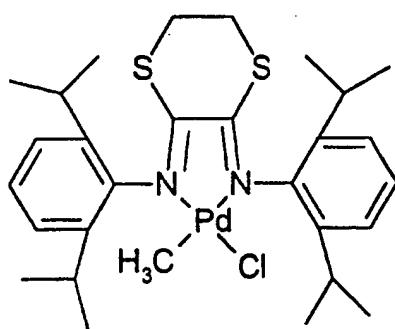
Preparation of the nickel dibromide complex of 1,4-dimethyl-2,3-bis(2,6-dimethylphenylimino)piperazine.

A 50 mL Schlenk flask equipped with a magnetic stir bar and capped with a septum was charged with 48 mg of 1,4-dimethyl-2,3-bis(2,6-dimethylphenylimino)piperazine and 35 mg of (1,2-dimethoxyethane)nickel(II) dibromide under an inert atmosphere. Dry, deoxygenated dichloromethane (5 mL) was added and the mixture was stirred under an argon atmosphere, turning green within about 5 minutes and slowly producing a green precipitate. After a total of 7 hours, the volatiles were removed under reduced pressure (1 mm Hg) and the residue was washed with 2 x 5 mL of dry, deoxygenated diethyl ether. The resultant green solid was dried under reduced pressure (1 mm Hg).

15

Example 34

20 Synthesis of:



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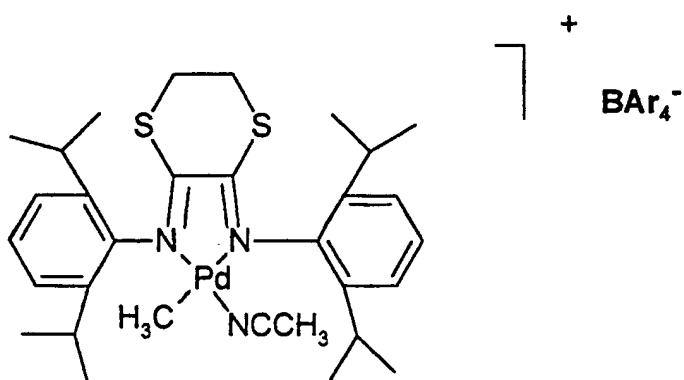
In the glove box, a Schlenk flask was charged with 500 mg of 2,3-bis(2,6-diisopropylphenylimino)-[1,4]dithiane and 250 mg (1,5-cyclooctadiene)palladium methyl chloride. The flask was removed from the box and placed under an argon atmosphere. To the solid mixture was 5 added 20 ml of methylene chloride resulting in an orange solution. The mixture was left to stir for 4 hours. After 4 hours, 20 ml of hexane was added resulting in the precipitation of an orange solid. The solvent was removed via filter cannula leaving a red/orange solid. The solid was subsequently washed 3 x 10 ml of hexane and dried *in vacuo* resulting in 10 490 mg of the complex (83 % yield). ^1H NMR is consistent with the proposed structure.

15

Example 35

20 Synthesis of:

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XXXV

In the glove box, a Schlenk flask was charged with 490 mg of 2,3-bis(2,6-diisopropylphenylimino)-[1,4]dithiane palladium methyl chloride and 5 738 mg of NaBAr₄ where Ar = 3,5-bis-trifluoromethylphenyl. The flask was removed from the box and placed under an argon atmosphere. To the solid mixture was added 25 ml of methylene chloride and 0.2 ml of acetonitrile resulting in an orange solution. The mixture was left to stir for 3 hours. After 3 hours, the solution was transferred via filter cannula leaving a gray 10 solid (NaCl). The solvent was subsequently removed *in vacuo* resulting in a orange glass (1.1 g of the complex, 90 % yield). ¹H NMR is consistent with the proposed structure.

15

Example 36

Polymerization of ethylene with the nickel dibromide complex of 2,3-bis(2,6-dimethylphenylimino)-[1,4]dithiane in the presence of MAO.

A 200 mL pear-shaped Schlenk flask equipped with a magnetic stir bar and capped with a septum was charged with 5.3 mg of the nickel dibromide complex of 2,3-bis(2,6-dimethylphenylimino)-[1,4]dithiane. The 20 flask was evacuated and refilled with ethylene, then charged with 75 mL of

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dry, deoxygenated toluene. The resultant suspension was cooled to 0 °C and allowed to equilibrate with 1 atmosphere ethylene for 15 minutes, then treated with 4.0 mL of a 10 wt% solution of MAO in toluene and stirred under 1 atmosphere ethylene. A white polyethylene precipitate (with a faint red-brown tinge) was observed within minutes. After 10 minutes, the mixture was quenched by the addition of acetone (50 mL), methanol (50 mL) and 6 N aqueous HCl (100 mL). The swollen polyethylene which separated was isolated by vacuum filtration and washed with water, methanol and acetone, then dried under reduced pressure (0.05-0.1 mm Hg) for 48 hours to give 2.5 g of a white polyethylene. A similar reaction at 21.5° C using 0.104 mg of the nickel complex (100 µL of a 1.04 mg/mL stock solution in *o*-difluorobenzene) gave 426 mg polyethylene after 14 minutes reaction (359,000 TO/h). ¹H NMR: 24 branches/1000 carbon atoms. GPC: $M_n = 810,000$; $M_w/M_n = 2.3$.

15

Example 37

Polymerization of ethylene using a catalyst generated *in situ* from 2,3-bis(2,6-dimethylphenylimino)-[1,4]dithiane, bis(1,5-cyclooctadiene)nickel(0) and HB(Ar)₄ (Ar = 3,5-bis(trifluoromethyl)phenyl).

20 A 250 mL pear-shaped Schlenk flask equipped with a magnetic stir bar and capped with a septum was charged with 20 mg of bis(1,5-cyclooctadiene)nickel(0), 33 mg of 2,3-bis(2,6-dimethylphenylimino)-[1,4]dithiane, and 83 mg of the ether solvate HB(Ar)₄. The flask was evacuated and refilled with ethylene, then charged with 75 mL of dry, deoxygenated toluene. The deep violet solution which resulted was stirred under ethylene at 25 °C for 30 minutes, then quenched by addition of acetone (50 mL), and methanol (50 mL). The polyethylene which separated was isolated by vacuum filtration and washed with acetone, then dried under reduced pressure (0.5 mm Hg) for 18 hours to give 339 mg of

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white polyethylene (332 TO/h). ^1H NMR: 47 branches/1000 carbon atoms. GPC: $M_n = 180,000$; $M_w/M_n = 2.4$.

Example 38

5 Polymerization of ethylene with the nickel dibromide complex of 2,3-bis(2,6-dimethylphenylimino)-[1,4]dithiane in the presence of MAO.
A 200 mL pear-shaped Schlenk flask equipped with a magnetic stir bar was charged with 0.5 mL of a stock solution (10 mg in 10 mL CH_2Cl_2) of the nickel dibromide complex of 2,3-bis(2,6-

10 dimethylphenylimino)-[1,4]dithiane. The flask was evacuated and refilled with ethylene, and charged with 75 mL of dry, deoxygenated toluene. The reaction flask was placed in a water bath (23° C) and treated with 1.0 mL of a 10 wt% solution of MAO in toluene and stirred under 1 atmosphere ethylene. A white polyethylene precipitate was observed within seconds.

15 After 5 minutes, the mixture was quenched by the addition of acetone, methanol and 6 N aqueous HCl. The swollen polyethylene which separated was isolated by vacuum filtration and washed with acetone. The resulting polymer was dried for several hours in a vacuum oven at 80° C. 580 mg of a white rubbery solid was isolated (285,000 TO/h). DSC: (2nd heat) broad

20 melt with an endothermic maximum at 87° C. ^1H NMR, 37 branches/1000 carbon atoms. GPC: $M_n = 186,000$; $M_w/M_n = 2.06$.

Example 39

25 Polymerization of ethylene with the nickel dibromide complex of 2,3-bis(2,6-diisopropylphenylimino)-[1,4]dithiane in the presence of MAO.
A 200 mL pear-shaped Schlenk flask equipped with a magnetic stir bar was charged with 0.5 mL of a stock solution (10 mg in 10 mL CH_2Cl_2) of the nickel dibromide complex of 2,3-bis(2,6-diisopropylphenylimino)-[1,4]dithiane. The flask was evacuated and refilled with ethylene, and charged with 75 mL of dry, deoxygenated toluene. The

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reaction flask was placed in a water bath and treated with 1.0 mL of a 10 wt% solution of MAO in toluene and stirred under 1 atmosphere ethylene. After 10 minutes, the mixture was quenched by the addition of acetone, methanol and 6 N aqueous HCl. The swollen polyethylene which separated 5 was isolated by vacuum filtration and washed with acetone. The resulting polymer was dried for several hours in a vacuum oven at 80° C. 210 mg of a white rubbery amorphous polymer was isolated (63,000 TO/h). DSC: (2nd heat) broad melt with an endothermic maximum at 6 °C. ^1H NMR: 92 branches/1000 carbon atoms. GPC: $M_n = 146,000$; $M_w/M_n = 1.85$.

10

Example 40

Polymerization of ethylene with the nickel dibromide complex of 2,3-bis(2,6-dimethylphenylimino)-[1,4]dithiane in the presence of MMAO (modified methylalumoxane; 23 % iso-butylaluminoxane).

15 A 600 mL Parr® autoclave was first heated to about 100° C under high vacuum to ensure that the reactor was dry. The reactor was cooled and purged with argon. Under an argon atmosphere, the autoclave was charged with 150 mL of toluene and 0.3 mL of a stock solution (10 mg in 10 mL CH_2Cl_2) of the nickel dibromide complex of 2,3-bis(2,6-dimethylphenylimino)-[1,4]dithiane. The autoclave was heated to 40° C and 20 2 mL of MMAO in heptane (6.42 wt% aluminum) was added. The reactor was rapidly pressurized to 100 psig with ethylene and the temperature ramped up to 50° C. After 10 minutes at 50° C, the reaction was quenched by the addition of acetone, and methanol. The swollen polyethylene which 25 separated was isolated and dried for several hours in a vacuum oven at 80° C. 4.8 g of a white rubbery solid was isolated (2,000,000 TO/h). DSC: (2nd heat) broad melt with an endothermic maximum at 97 °C. ^1H NMR: 28 branches/1000 carbon atoms. GPC: $M_n = 155,000$; $M_w/M_n = 2.10$.

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Example 41

Polymerization of ethylene with the nickel dibromide complex of 2,3-bis(2,6-dimethylphenylimino)-[1,4]dithiane in the presence of MMAO (23 % isobutylaluminoxane).

5 A 600 mL Parr® autoclave was first heated to about 100° C under high vacuum to ensure that the reactor was dry. The reactor was cooled and purged with argon. Under an argon atmosphere, the autoclave was charged with 150 mL of toluene and 0.3 mL of a stock solution (10 mg in 10 mL CH₂Cl₂) of the nickel dibromide complex of 2,3-bis(2,6-dimethylphenylimino)-[1,4]dithiane. The autoclave was cooled to 15° C and 10 2 mL of MMAO in heptane (6.42 wt% aluminum) was added. The reactor was rapidly pressurized to 100 psig with ethylene and the temperature ramped up to 25° C. After 10 minutes at 25° C, the reaction was quenched by the addition of acetone, and methanol. The swollen polyethylene which 15 separated was isolated and dried for several hours in a vacuum oven at 80° C. 4.4 g of a white rubbery polyethylene was isolated (1,800,000 TO/h). DSC: (2nd heat) melt with an endothermic maximum at 125 °C. ¹H NMR: 20 6 branches/1000 carbon atoms. GPC: M_n = 598,000; M_w/M_n = 2.12.

Example 42

Polymerization of ethylene with the nickel dibromide complex of 2,3-bis(2,6-dimethylphenylimino)-[1,4]dithiane in the presence of MMAO (23 % isobutylaluminoxane).

25 A 600 mL Parr® autoclave was first heated to about 100° C under high vacuum to ensure that the reactor was dry. The reactor was cooled and purged with argon. Under an argon atmosphere, the autoclave was charged with 150 mL of toluene and 1.0 mL of a stock solution (10 mg in 10 mL CH₂Cl₂) of the nickel dibromide complex of 2,3-bis(2,6-dimethylphenylimino)-[1,4]dithiane. The autoclave was heated to 55° C and 30 2 mL of MMAO in heptane (6.42 wt% aluminum) was added. The reactor

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was rapidly pressurized to 100 psig with ethylene and the temperature ramped up to 65° C. After 10 minutes at 65° C, the reaction was quenched by the addition of acetone, and methanol. The swollen polyethylene which separated was isolated and dried for several hours in a vacuum oven at 80°
5 C. 5.3 g of a white rubbery solid was isolated (640,000 TO/h). DSC: (2nd heat) melt with an endothermic maximum at 78° C. ¹H NMR: 47 branches/1000 carbon atoms. GPC: $M_n = 86,000$; $M_w/M_n = 1.95$.

Example 43

10 Polymerization of ethylene with the nickel dibromide complex of 2,3-bis(2,6-dimethylphenylimino)-[1,4]dithiane in the presence of MMAO (23 % isobutylalumininoxane).

A 600 mL Parr® autoclave was first heated to about 100° C under high vacuum to ensure that the reactor was dry. The reactor was cooled
15 and purged with argon. Under an argon atmosphere, the autoclave was charged with 150 mL of toluene and 1.0 mL of a stock solution (10 mg in 10 mL CH₂Cl₂) of the nickel dibromide complex of 2,3-bis(2,6-dimethylphenylimino)-[1,4]dithiane. The autoclave was heated to 70° C and 2 mL of MMAO in heptane (6.42 wt% aluminum) was added. The reactor
20 was rapidly pressurized to 100 psig with ethylene and the temperature ramped up to 80° C. After 10 minutes at 80° C, the reaction was quenched by the addition of acetone, and methanol. The swollen polyethylene which separated was isolated and dried for several hours in a vacuum oven at 80° C. 3.5 g of a white rubbery solid was isolated (440,000 TO/h). DSC: (2nd heat) melt with an endothermic maximum at 67° C. ¹H NMR: 53 branches/1000 carbon atoms. GPC: $M_n = 87,000$; $M_w/M_n = 1.66$.

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Example 44

Polymerization of ethylene with the nickel dibromide complex of 2,3-bis(2,6-diisopropylphenylimino)-[1,4]dithiane in the presence of MMAO (23 % iso-butylalumininoxane).

5 A 600 mL Parr® autoclave was first heated to about 100° C under high vacuum to ensure that the reactor was dry. The reactor was cooled and purged with argon. Under an argon atmosphere, the autoclave was charged with 150 mL of toluene and 0.5 mL of a stock solution (10 mg in 10 mL CH₂Cl₂) of the nickel dibromide complex of 2,3-bis(2,6-di-isopropylphenylimino)-[1,4]dithiane. The autoclave was heated to 40° C and 2 mL of MMAO in heptane (6.42 wt% aluminum) was added. The reactor was rapidly pressurized to 100 psig with ethylene and the temperature ramped up to 50° C. After 10 minutes at 50° C, the reaction was quenched by the addition of acetone, and methanol. The swollen polyethylene which separated was isolated and dried for several hours in a vacuum oven at 80° C. 2.4 g of a white rubbery solid was isolated (700,000 TO/h). DSC: (2nd heat) melt with an endothermic maximum at 46° C. ¹H NMR: 75 branches/1000 carbon atoms. GPC: M_n = 966,000; M_w/M_n = 1.70.

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Example 45

Polymerization of ethylene with the nickel dibromide complex of 2,3-bis(2,6-diisopropylphenylimino)-[1,4]dithiane in the presence of MMAO (23 % iso-butylalumininoxane).

25 The procedure described in example 44 was followed except the polymerization was conducted at 80 °C. 1.4 g of a white rubbery solid was isolated (400,000 TO/h). DSC: (2nd heat) melt with an endothermic maximum at 0° C. ¹H NMR: 95 branches/1000 carbon atoms. GPC: M_n = 406,000; M_w/M_n = 2.05.

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Example 46

Polymerization of ethylene with the nickel dibromide complex of 2,3-bis(2,6-diisopropylphenylimino)-[1,4]dithiane in the presence of MMAO (23 % iso-butylalumininoxane).

5 The procedure described in Example 44 was followed except the polymerization was conducted at 65° C. 2.15 g of a white rubbery solid was isolated (630,000 TO/h). DSC: (2nd heat) melt with an endothermic maximum at 15° C. ^1H NMR: 89 branches/1000 carbon atoms. GPC: $M_n = 502,000$; $M_w/M_n = 1.78$.

10

Example 47

Polymerization of ethylene with the nickel dibromide complex of 2,3-bis(2,6-diisopropylphenylimino)-[1,4]dithiane in the presence of MMAO(23 % iso-butylalumininoxane).

15

The procedure described in Example 44 was followed except the polymerization was conducted at 25 ° C. 1.9 g of a white rubbery solid was isolated (560,000 TO/h). DSC: (2nd heat) melt with an endothermic maximum at 90 ° C. ^1H NMR: 33 branches/1000 carbon atoms. GPC: $M_n = 839,000$; $M_w/M_n = 1.37$.

20

Example 48

Oligomerization of ethylene to α -olefin with the nickel dibromide complex of 2,3-bis(phenylimino)-[1,4]dithiane in the presence of MAO.

25 A 1 L Fischer-Porter bottle was assembled onto a pressure head equipped with a mechanical stirrer and gas and liquid feed-through ports, then pressurized to 75 psig of ethylene and relieved to ambient pressure seven times. The bottle was immersed in a 21.5° C. water bath, then 50 mL of dry, deoxygenated toluene was added via syringe, followed by 100 μL of

30

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a stock solution of 15.3 mg of the nickel dibromide complex of 2,3-bis(phenylimino)-[1,4]dithiane in 15.0 mL of dichloromethane, followed by another 50 mL of toluene. The mixture was stirred at 300 rpm under 75 psig ethylene for 5 minutes to saturate the solution with ethylene, then the 5 pressure was relieved, and 4.0 mL of a 10 wt% solution of MAO in toluene was quickly added. The flask was immediately re-pressurized to 75 psig ethylene and stirred at 300 rpm. After 30 minutes, the pressure was relieved and the reaction quenched by addition of 10 mL of methanol. After disassembling the apparatus, another 40 mL of methanol, 50 mL of 6 N 10 aqueous HCl, and 10 mL acetone were added and the mixture was stirred to complete hydrolysis of the MAO. The resultant organic layer was separated, washed with 6 N aqueous HCl (1 x 25 mL), and water (2 x 50 mL), then concentrated under reduced pressure (15 Torr) at 40° C to obtain an oil. This was treated with toluene (50 mL) and re-concentrated twice, 15 then treated with acetone (50 mL) and re-concentrated, to obtain a waxy white polyethylene solid. Drying *in vacuo* at 100° C, 250 mm Hg for 14 hours gave 0.180 g of polymer, approximate M_n = 517, containing approximately 85% α -olefin and 15% internal olefin.

20

Example 49

Polymerization of ethylene with the nickel dibromide complex of 2,3-bis(2,6-dimethylphenylimino)-2,3-dihydrobenzo[1,4]dithiine in the presence of MAO.

25 A 200 mL pear-shaped Schlenk flask equipped with a magnetic stir bar and capped with a septum was charged with 100 mL of dry, deoxygenated toluene. The flask was placed in a water bath and allowed to equilibrate with 1 atmosphere ethylene for 10 minutes, then 0.25 mL of a stock solution prepared from 10.2 mg of the nickel dibromide complex of 30 2,3-bis-(2,6-dimethylphenylimino)-2,3-dihydrobenzo[1,4]dithiine in 13.11 g

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dry, deoxygenated dichloromethane was added. The reaction mixture was then treated with 4.0 mL of a 10 wt% solution of MAO in toluene and stirred under 1 atmosphere ethylene. Ethylene uptake and formation of a polyethylene precipitate were observed. After 6.5 minutes, the mixture was 5 quenched by the addition of acetone (50 mL), methanol (50 mL) and 6 N aqueous HCl (100 mL). The swollen polyethylene which separated was isolated by vacuum filtration, then dried at 80° C *in vacuo* for several hours. 287 mg of a white powdery polyethylene was isolated (236,000 TO/h). DSC: (2nd heat) melt with an endothermic maximum at 88° C. ¹H NMR 10 showed this material to contain approximately 36 branches/1000 carbon atoms. GPC: $M_n = 145,000$; $M_w/M_n = 2.35$.

Example 50

15 Polymerization of ethylene using a catalyst formed in situ from 2,3-bis(2,6-diisopropylphenylimino)-1,4-dithiane and [Pd(NCCH₃)₄][BF₄]₂

A 200 mL pear-shaped Schlenk flask equipped with a magnetic stir bar and capped with a septum was charged with 0.022 g of 2,3-bis(2,6-diisopropylphenylimino)-1,4-dithiane and 0.019 g [Pd(NCCH₃)₄][BF₄]₂. The flask was evacuated and refilled with ethylene, then 100 mL of dry, 20 deoxygenated dichloromethane was added via syringe and the resultant mixture was stirred under 1 atmosphere of ethylene at 25° C. Very little ethylene uptake was observed. After 10 minutes, 0.412 g B(C₆F₅)₃ was added, resulting in an increased rate of ethylene uptake. After a total of 84 minutes, the reaction was worked up by evaporating the dichloromethane 25 under a stream of nitrogen, washing the residue with methanol repeatedly to extract the B(C₆F₅)₃, and drying the residue *in vacuo* to obtain 0.57 g of amorphous polyethylene, approximate $M_n = 17,000$; $M_w/M_n = 1.3$. ¹H NMR showed approximately 105 branches per 1000 carbons.

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Example 51

Polymerization of ethylene with the nickel dibromide complex of 2,3- bis(4-methoxy-2,6-dimethylphenylimino)-[1,4]dithiane in the presence of MAO.

A 500 mL round bottom flask fitted with a Schlenk adapter and 5 equipped with a magnetic stir bar and capped with a septum was evacuated, flame-dried, then refilled with ethylene. The flask was provided with a room temperature (ca. 23°C) water bath, then charged with 100 mL of dry, deoxygenated toluene and allowed to equilibrate with 1 atmosphere ethylene for 30 minutes. The reaction mixture was then treated with 4.0 mL 10 of a 10 wt% solution of MAO in toluene and stirred under 1 atmosphere ethylene, then 0.10 mL of a stock solution (prepared from 6.3 mg of the nickel dibromide complex of 2,3- bis(4-methoxy-2,6-dimethylphenylimino)-[1,4]dithiane and 6.5 mL dichloromethane) was added. After 10 minutes, 15 the reaction mixture was quenched by the addition of acetone, methanol and 6 N aqueous HCl. The polyethylene which separated was isolated by vacuum filtration and washed with water, methanol and acetone, dried on the filter for 2 h, then further dried 13 days in a vacuum oven at 80°C. 182 mg of a white polyethylene was isolated (254,000 TO/h). DSC: (2nd heat) 20 melt with an endothermic maximum at 124° C. ¹H NMR: 13 branches/1000 carbon atoms. GPC: $M_n = 145,600$; $M_w/M_n = 2.6$.

Example 52

Polymerization of ethylene with the nickel dibromide complex of 2,3-bis(2,6-diisopropylphenylimino)-[1,4]dithiane in the presence of (CH₃CH₂)₂AlCl.

25 A 600 mL Parr [®] autoclave was first heated to about 100 °C under high vacuum to ensure the reactor was dry. The reactor was cooled and purged with argon. Under an argon atmosphere, the autoclave was charged with 150 mL of toluene and 0.5 mL of a stock solution (10 mg in 10 mL CH₂Cl₂) of the nickel dibromide complex of 2,3-bis(2,6- 30 diisopropylphenylimino)-[1,4]dithiane. The autoclave was heated to 45 °C

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and 2 mL of $(CH_3CH_2)_2AlCl$ (5000 equiv.) in toluene was added. The reactor was rapidly pressurized to 100 psig and the temperature ramped up to 50 °C. After 10 minutes at 50 °C, the reaction was quenched by the addition of acetone, and methanol. The swollen polyethylene which 5 separated was isolated by filtration and dried for several hours in a vacuum oven at 80 °C resulting in 1.9 g of a white rubbery solid (560,000 TO/h). DSC: (2nd heat) broad melt with an endothermic maximum at 30 °C. 1H NMR: 87 branches/1000 carbon atoms. GPC: Mn = 557,000; Mw/Mn = 1.82.

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Example 53

Polymerization of ethylene with the nickel dibromide complex of 2,3-bis(2,6-diisopropylphenylimino)-[1,4]dithiane in the presence of $(CH_3CH_2)_2AlCl$

A 600 mL Parr ® autoclave was first heated to about 100 °C under 15 high vacuum to ensure the reactor was dry. The reactor was cooled and purged with argon. Under an argon atmosphere, the autoclave was charged with 150 mL of toluene and 0.5 mL of a stock solution (10 mg in 10 mL CH_2Cl_2) of the nickel dibromide complex of 2,3-bis(2,6-diisopropylphenylimino)-[1,4]dithiane. The autoclave was heated to 45 °C 20 and 0.2 mL of $(CH_3CH_2)_2AlCl$ (500 equiv.) in toluene was added. The reactor was rapidly pressurized to 100 psig and the temperature ramped up to 50 °C. After 10 minutes at 50 °C, the reaction was quenched by the addition of acetone, and methanol. The swollen polyethylene which 25 separated was isolated by filtration and dried for several hours in a vacuum oven at 80 °C resulting in 2 g of a white rubbery solid (590,000 TO/h). DSC: (2nd heat) broad melt with an endothermic maximum at 30 °C. 1H NMR: 85 branches/1000 carbon atoms. GPC: Mn = 515,000; Mw/Mn = 1.81.

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Example 54

Polymerization of ethylene with the nickel dibromide complex of 2,3-bis(2,6-diisopropylphenylimino)-[1,4]dithiane in the presence of (CH₃CH₂)₂AlCl

A 600 mL Parr[®] autoclave was first heated to about 100 °C under 5 high vacuum to ensure the reactor was dry. The reactor was cooled and purged with argon. Under an argon atmosphere, the autoclave was charged with 150 mL of toluene and 0.5 mL of a stock solution (10 mg in 10 mL CH₂Cl₂) of the nickel dibromide complex of 2,3-bis(2,6-diisopropylphenylimino)-[1,4]dithiane. The autoclave was heated to 45 °C 10 and 0.04 mL of (CH₃CH₂)₂AlCl (100 equiv.) in toluene was added. The reactor was rapidly pressurized to 100 psig and the temperature ramped up to 50 °C. After 10 minutes at 50 °C, the reaction was quenched by the addition of acetone, and methanol. The swollen polyethylene which 15 separated was isolated by filtration and dried for several hours in a vacuum oven at 80 °C resulting in 1.5 g of a white rubbery solid (440,000 TO/h). DSC: (2nd heat) broad melt with an endothermic maximum at 29 °C. ¹H NMR: 80 branches/1000 carbon atoms. GPC: Mn = 422,000; Mw/Mn = 1.97.

20

Example 55

Copolymerization of ethylene and ethyl undecenoate with the nickel dibromide complex of 2,3-bis(2,6-dimethylphenylimino)-[1,4]dithiane in the presence of MMAO (23 % iso-butylalumininoxane).

A flame dried Schlenk flask equipped with a stir bar and a rubber 25 septum was charged with 50 ml of toluene and 5 mg of the nickel dibromide complex of 2,3-bis(2,6-dimethylphenylimino)-[1,4]dithiane. The flask was cooled to 0 °C in an ice-water bath and filled with ethylene (1 atmosphere). To the flask was added 2.0 ml of MMAO in heptane (6.42 wt% aluminum). Within 5 seconds, 5 ml of ethyl undecenoate was added to give a purple 30 solution. The mixture was left to stir for 16 hours. Acetone, methanol and

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6M HCl were added to quench the reaction and precipitate the polymer. The polymer was collected by suction filtration and washed with copious amounts of acetone to ensure all of the ethyl undecenoate comonomer was removed resulting in 100 mg of white powdery polymer. NMR spectroscopic analysis is consistent with the preparation of an ester group containing copolymer. In addition, ethylene homopolymer, which resulted from the short reaction time prior to addition of the ethyl undecenoate, was present. ^1H NMR: 7.5 wt % ethyl undecenoate incorporated. GPC: $M_n = 9500$, $M_w/M_n = 16.6$. DSC: $T_m = 128^\circ\text{C}$.

10

Example 56

Co-polymerization of ethylene and 1,13-tetradecadiene with the nickel dibromide complex of 2,3-bis(2,6-dimethylphenylimino)-[1,4]dithiane in the presence of MAO.

15 A 200 mL pear-shaped Schlenk flask equipped with a magnetic stir bar and capped with a septum was flame-dried under vacuum, refilled with ethylene, and then sequentially charged with 50 mL of dry, deoxygenated toluene, 6.0 mL of deoxygenated 1,13-tetradecadiene, and 1.0 mL of a stock solution of 11.8 mg of the nickel dibromide complex of 2,3-bis(2,6-dimethylphenylimino)-[1,4]dithiane in 10.0 mL of dry, deoxygenated dichloromethane. The flask was placed in a 23°C water bath and allowed to equilibrate with 1 atmosphere of ethylene for 5 minutes, then 4.0 mL of a 10 wt% solution of MAO in toluene was added and the mixture was stirred under 1 atmosphere of ethylene. Ethylene uptake was observed and the mixture rapidly became more viscous. After 7 minutes, the reaction was quenched by the addition of acetone (50 mL), methanol (50 mL) and 6 N aqueous HCl (100 mL). The co-polymer which separated was isolated by vacuum filtration and dried in *vacuo* at 100°C for 24 hours to obtain 0.72 g of a rubbery white polymer, which formed a gel upon attempted re-dissolution in hot o-dichlorobenzene.

Example 57Polymerization of ethylene with the nickel dibromide complex of 2,3-bis(2,6-dimethylphenylimino)-[1,4]dioxane in the presence of MAO.

5 A 200 mL pear-shaped Schlenk flask equipped with a magnetic stir bar and capped with a septum was charged with 3.4 mg of the nickel dibromide complex of 2,3-bis(2,6-dimethylphenylimino)-[1,4]dioxane. The flask was evacuated and refilled with ethylene, then charged with 75 mL of dry, deoxygenated toluene. The resultant suspension was cooled to 0° C
10 and allowed to equilibrate with 1 atmosphere ethylene for 15 minutes, then treated with 4.0 mL of a 10 wt% solution of MAO in toluene and stirred under 1 atmosphere ethylene. A white polyethylene precipitate (with a faint yellow-orange tinge) was observed within minutes. After 38 minutes, the mixture was quenched by the addition of acetone (50 mL), methanol (50 mL) and 6 N aqueous HCl (100 mL). The swollen polyethylene that separated was isolated by vacuum filtration and washed with water, methanol and acetone, then dried under reduced pressure (0.05-0.1 mm Hg) for 18 hours to give 6.0 g of a white polyethylene (54,000 TO/h). ¹H NMR: 19 branches/1000 carbon atoms. GPC: M_n = 504,000; M_w/M_n = 2.3.

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Example 58Polymerization of ethylene with the nickel dibromide complex of 2,3-bis(benzylloxymethyl)-5,6-bis(2,6-dimethylphenylimino)-[1,4]dioxane in the presence of MAO.

25 A 500 mL round bottom flask was fitted with a Schlenk adapter and equipped with a magnetic stir bar and capped with a septum was charged with 100 mL of dry, deoxygenated toluene. The flask was placed in a water bath and allowed to equilibrate with 1 atmosphere ethylene for 19 minutes, then 0.25 mL of a stock solution prepared from 10.0 mg of the nickel
30 dibromide complex of 2,3-bis(benzylloxymethyl)-5,6-bis(2,6-

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dimethylphenylimino)-[1,4]dioxane and 10.0 mL dichloromethane was added. The reaction mixture was then treated with 4.0 mL of a 10 wt% solution of MAO in toluene and stirred under 1 atmosphere ethylene. Ethylene uptake and formation of a polyethylene precipitate were observed.

5 After 6.5 minutes, the mixture was quenched by the addition of acetone, methanol and 6 N aqueous HCl. The swollen polyethylene which separated was isolated by vacuum filtration, then dried at 80° C *in vacuo* for several hours. 392 mg of white polyethylene was isolated (404,000TO/h). DSC: (2nd heat) melt with an endothermic maximum at 120 C. ¹H NMR: 16

10 branches/1000 carbon atoms. GPC: $M_n = 125,000$; $M_w/M_n = 2.8$.

Example 59

Polymerization of ethylene with the nickel dibromide complex of 5-methoxymethyl-2,3-bis(2,6-dimethylphenylimino)-[1,4]dioxane in the presence of MAO.

A 200 mL pear-shaped Schlenk flask equipped with a magnetic stir bar and capped with a septum was charged with 3.8 mg of the nickel dibromide complex of 5-methoxymethyl-2,3-bis(2,6-dimethylphenylimino)-[1,4]dioxane. The flask was evacuated and refilled with ethylene, then charged with 75 mL of dry, deoxygenated toluene. The resultant suspension was cooled to 0 °C. and allowed to equilibrate with 1 atmosphere ethylene for 15 minutes, then treated with 4.0 mL of a 10 wt% solution of MAO in toluene and stirred under 1 atmosphere ethylene. After 10 minutes, the mixture was quenched by the addition of acetone (50 mL),

20 methanol (50 mL) and 6 N aqueous HCl (100 mL). The swollen polyethylene which separated was isolated by vacuum filtration and washed with water, methanol and acetone, then dried under reduced pressure (0.05-0.1 mm Hg) for 48 hours to give 1.04 g of a white, powdery polyethylene. A similar reaction, also at 0° C, was conducted using 0.655 g

25 (equivalent to 0.57 mg of the nickel complex) of a stock solution of 11.6 mg

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nickel dibromide complex of 5-methoxymethyl-2,3-bis-(2,6-dimethylphenylimino)-[1,4]dioxane in 13.238 g dichloromethane to obtain 291 mg white, powdery polyethylene after 15 minutes reaction.

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Example 60

Polymerization of ethylene with the nickel dibromide complex of 2,3-bis(2,6-dimethylphenylimino)-[1,4]dioxane in the presence of MMAO (23 % isobutylalumininoxane).

A 600 mL Parr [®] autoclave was first heated to about 100 °C under high vacuum to ensure the reactor was dry. The reactor was cooled and purged with argon. Under an argon atmosphere, the autoclave was charged with 150 mL of toluene and 0.3 mL of a stock solution (10 mg in 10 mL CH₂Cl₂) of the nickel dibromide complex of 2,3-bis(2,6-dimethylphenylimino)-[1,4]dioxane. The autoclave was heated to 60 °C and 10 2 mL of MMAO in heptane (6.42 wt% aluminum) was added. The reactor was rapidly pressurized to 100 psig and the temperature ramped up to 65 °C. After 10 minutes at 65 °C, the reaction was quenched by the addition of acetone, and methanol. The swollen polyethylene which separated was 15 isolated by filtration and dried for several hours in a vacuum oven at 80° C. 20 1.3 g of a white rubbery solid was isolated (480,000 TO/h). DSC: (2nd heat) melt with an endothermic maximum at 76 °C. ¹H NMR: 33 branches/1000 carbon atoms. GPC: Mn = 42,000; Mw/Mn = 1.82.

Example 61

25 Polymerization of ethylene with the nickel dibromide complex of 2,3-bis(2,6-dimethylphenylimino)-[1,4]dioxane in the presence of MMAO (23 % isobutylalumininoxane).

The procedure described in example 60 was followed except the polymerization was conducted at 25 °C resulting in 0.59 g of polyethylene 30 (226,000 TO/h). DSC: (2nd heat) melt with an endothermic maximum at

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125 °C. ^1H NMR: 9 branches/1000 carbon atoms. GPC: Mn = 237,000;
Mw/Mn = 2.15.

Example 62

5 Polymerization of ethylene with the nickel dibromide complex of 2,3-bis(2,6-dimethylphenylimino)-[1,4]dioxane in the presence of MMAO (23 % iso-butylalumininoxane).

The procedure described in example 60 was followed except the polymerization was conducted at 80 °C resulting in 0.29 g of polyethylene
10 (110,000 TO/h). ^1H NMR: 69 branches/1000 carbon atoms. GPC: Mn = 23,000; Mw/Mn = 1.65.

Example 63

15 Polymerization of ethylene with the nickel dibromide complex of 2,3-bis(2,6-dimethylphenylimino)-[1,4]dioxane in the presence of MMAO (23 % iso-butylalumininoxane).

The procedure described in example 60 was followed except the polymerization was conducted at 50 °C resulting in 3.1 g of polyethylene
20 (1,200,000 TO/h). DSC: (2nd heat) melt with an endothermic maximum at 95 °C. ^1H NMR: 48 branches/1000 carbon atoms. GPC: Mn = 63,000;
Mw/Mn = 1.92.

Example 64

25 Polymerization of ethylene with the nickel dibromide complex of 2,3-bis(2,6-diisopropylphenylimino)-[1,4]dioxane in the presence of MMAO (23 % iso-butylalumininoxane).

A 600 mL Parr [®] autoclave was first heated to about 100 °C under high vacuum to ensure the reactor was dry. The reactor was cooled and purged with argon. Under an argon atmosphere, the autoclave was charged
30 with 150 mL of toluene and 0.5 mL of a stock solution (10 mg in 10 mL

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CH₂Cl₂) of the nickel dibromide complex of 2,3-bis(2,6-diisopropylphenylimino)-[1,4]dioxane. The autoclave was heated to 60 °C and 2 mL of MMAO in heptane (6.42 wt% aluminum) was added. The reactor was rapidly pressurized to 100 psig and the temperature ramped up 5 to 65 °C. After 10 minutes at 65 °C, the reaction was quenched by the addition of acetone, and methanol. The swollen polyethylene which separated was isolated by filtration and dried for several hours in a vacuum oven at 80 °C. 2.5 g of a white rubbery solid was isolated (660,000 TO/h). DSC: (2nd heat) broad melt with an endothermic maximum at 30 °C. ¹H 10 NMR: 82 branches/1000 carbon atoms. GPC: Mn = 147,000; Mw/Mn = 1.91.

Example 65

Polymerization of ethylene with the nickel dibromide complex of 2,3-bis(2,6-diisopropylphenylimino)-[1,4]dioxane in the presence of MMAO (23 % iso-butylaluminoxane).

The procedure described in example 64 was followed except the polymerization was conducted at 50 °C resulting in 3.4 g of polyethylene (900,000 TO/h). DSC: (2nd heat) melt with an endothermic maximum at 20 50 °C. ¹H NMR: 65 branches/1000 carbon atoms. GPC: Mn = 219,000; Mw/Mn = 1.85.

Example 66

Polymerization of ethylene with the nickel dibromide complex of 2,3-bis(2,6-diisopropylphenylimino)-[1,4]dioxane in the presence of MMAO (23 % iso-butylaluminoxane).

The procedure described in example 64 was followed except the polymerization was conducted at 25 °C resulting in 1.22 g of polyethylene (320,000 TO/h). DSC: (2nd heat) melt with an endothermic maximum at

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112 °C. ^1H NMR: 17 branches/1000 carbon atoms. GPC: Mn = 476,000; Mw/Mn = 2.02.

Example 67

5 Polymerization of ethylene with the nickel dibromide complex of 2,3-bis(2,6-diisopropylphenylimino)-[1,4]dioxane in the presence of MMAO (23 % iso-butylalumininoxane).

The procedure described in example 64 was followed except the polymerization was conducted at 80 °C resulting in 0.9 g of polyethylene 10 (240,000 TO/h). DSC: (2nd heat) melt with an endothermic maximum at -10 °C. ^1H NMR: 99 branches/1000 carbon atoms. GPC: Mn = 98,800; Mw/Mn = 1.81.

Example 68

15 Copolymerization of ethylene and ethyl undecenoate with the nickel dibromide complex of 2,3-bis(2,6-dimethylphenylimino)-[1,4]dioxane in the presence of MMAO (23 % iso-butylalumininoxane).

A flame dried Schlenk flask equipped with a stir bar and a rubber septum was charged with 50 ml of toluene and 6 mg of the nickel dibromide 20 complex of 2,3-bis(2,6-dimethylphenylimino)-[1,4]dioxane. The flask was cooled to 0 °C in an ice-water bath and filled with ethylene (1 atmosphere). To the flask was added, 2.0 ml of MMAO in heptane (6.42 wt% aluminum). Within 15 seconds 2.5 ml of ethyl undecenoate was added to give a purple 25 solution. The mixture was left to stir for 16 hours. Acetone, methanol and 6M HCl were added to quench the reaction and precipitate the polymer.

The polymer was collected by suction filtration and washed with copious amounts of acetone to ensure all of the ethyl undecenoate comonomer is removed resulting in 510 mg of white powdery polymer. NMR 30 spectroscopic analysis is consistent with the preparation of an ester group containing copolymer. In addition, ethylene homopolymer, which resulted

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from the short reaction time prior to addition of the ethyl undecenoate, was present. IR: CO stretch at 1742 cm⁻¹. ¹H NMR: 1.0 wt % ethyl undecenoate incorporated. GPC: M_n = 61,000, M_w/M_n = 6.4. DSC: T_m = 125 °C.

5

Example 69

Polymerization of ethylene with the nickel dibromide complex of 2,3-bis(2,6-diisopropylphenylimino)-4-methylmorpholine in the presence of MAO.

A 1 L Fischer-Porter bottle was assembled onto a pressure head equipped with a mechanical stirrer and gas and liquid feed-through ports, then 10 pressurized to 75 psig of ethylene and relieved to ambient pressure seven times. The bottle was immersed in a 54° C. water bath, then 100 mL of dry, deoxygenated toluene was added via syringe. The mixture was re-pressurized with ethylene at 75 psig and stirred at 300 rpm for 5 minutes to 15 saturate the solution with ethylene, then the pressure was again relieved, and 4.0 mL of a 10 wt% solution of MAO in toluene was quickly added. The apparatus was again re-pressured to 75 psig ethylene and stirred at 300 rpm for another 5 min to ensure saturation with ethylene. The pressure was once again relieved to ambient pressure and 0.5 mL of a stock solution 20 prepared from 10.0 mg of the nickel dibromide complex of 2,3-bis(2,6-diisopropylphenylimino)-4-methylmorpholine and 10 mL dichloromethane was quickly added and the system quickly pressurized once again with ethylene to 75 psig. After 7 min the pressure was relieved to atmospheric and the reaction was quenched by addition of 5 mL methanol. After the 25 apparatus was disassembled, an additional 50 mL methanol, 50 mL aqueous 6N HCl and 20 mL acetone was added. The resultant organic layer was separated, washed with 6 N aqueous HCl (1 x 25 mL), and water (2 x 50 mL), then concentrated by rotary evaporation under reduced pressure (10 Torr) at 40° C. The residue was then treated with toluene (50 30 mL) and re-concentrated to afford 134 mg of a very rubbery, clear

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polyethylene (55,000 TO/h). ^1H NMR: 134 branches/1000 carbon atoms.
GPC: $M_n = 169,000$; $M_w/M_n = 1.4$.

Example 70

5 Polymerization of ethylene with the nickel dibromide complex of 2,3-bis(2,6-dimethyl-phenylimino)-4-methylmorpholine in the presence of MMAO.

A 600 mL Parr $^{\circ}$ autoclave was first heated to about 100 $^{\circ}\text{C}$ under high vacuum to ensure the reactor was dry. The reactor was cooled and purged with argon. Under an argon atmosphere, the autoclave was charged 10 with 150 mL of toluene and 1 mL of a stock solution (10 mg in 20 mL CH_2Cl_2) of the nickel dibromide complex of 2,3-bis(2,6-dimethyl-phenylimino)-4-methylmorpholine. The autoclave was heated to 45 $^{\circ}\text{C}$ and 15 3 mL of MMAO in heptane (6.42 wt% aluminum) was added. The reactor was rapidly pressurized to 100 psig and the temperature ramped up to 50 $^{\circ}\text{C}$. After 10 minutes at 50 $^{\circ}\text{C}$, the reaction was quenched by the addition of acetone, and methanol. The swollen polyethylene which separated was isolated by filtration and dried for several hours in a vacuum oven at 80 $^{\circ}\text{C}$ resulting in 0.87 g of a white rubbery solid was isolated (138,000 TO/h). ^1H NMR: 97 branches/1000 carbon atoms.

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Example 71

Polymerization of ethylene with the nickel dibromide complex of 1,3-bis(4-methoxy-2,6-dimethylphenyl)-4,5-bis(4-methoxy-2,6-dimethylphenylimino)imidazolidin-2-one in the presence of MAO.

25 A 500 mL round bottom flask fitted with a Schlenk adapter and equipped with a magnetic stir bar and capped with a septum was evacuated, flame-dried, then refilled with ethylene. The flask was provided with a room temperature (ca. 23 $^{\circ}\text{C}$) water bath, then charged with 100 mL of dry, deoxygenated toluene and allowed to equilibrate with 1 atmosphere 30 ethylene for 30 minutes while stirring at 1000 rpm. The reaction mixture

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was then treated with 4.0 mL of a 10 wt% solution of MAO in toluene and stirred under 1 atmosphere ethylene, then 0.10 mL of a stock solution prepared from 6.0 mg of the nickel dibromide complex of 1,3-bis(4-methoxy-2,6-dimethylphenyl)-4,5-bis (4-methoxy-2,6-dimethylphenylimino)imidazolidin-2-one and 6.0 mL dichloromethane was added. After about 7 min and 20 seconds an additional 0.25 mL of the stock solution was added. After 15 more minutes, the reaction mixture was quenched by the addition of acetone, methanol and 6 N aqueous HCl. The polyethylene which separated was isolated by vacuum filtration and washed with water, methanol and acetone, then dried on the filter for 2 h, then further dried 13 days in a vacuum oven at 80°C to obtain 172 mg white polyethylene. DSC: (2nd heat) melt with an endothermic maximum at 124°C. ¹H NMR showed this material to contain approximately 18 branches/1000 carbon atoms. GPC: $M_n = 56,500$; $M_w/M_n = 3.55$.

15

Example 72

Polymerization of ethylene with the reaction product of 1,3-bis-(2,6-dimethyl-phenyl)-4,5-bis-(2,6-dimethyl-phenylimino)-imidazolidin-2-one, (1,2-dimethoxyethane)nickel(II) dibromide, and silver tetrafluoroborate in the presence of MAO.

A 500 mL round bottom flask fitted with a Schlenk adapter, capped with a septum, and equipped with a magnetic stir bar was charged with 100 mL of dry, deoxygenated toluene. The flask was placed in a water bath and allowed to equilibrate with 1 atmosphere ethylene for 10 minutes, then 0.10 mL of a stock solution (freshly prepared from 240 mg of the reaction product of 1,3-bis-(2,6dimethylphenyl)-4,5-bis-(2,6-dimethylphenylimino)-imidazolidin-2-one, (1,2-dimethoxyethane)nickel(II) dibromide, and silver tetrafluoroborate in 10 mL dry, deoxygenated dichloromethane) was added. The reaction mixture was then treated with 4.0 mL of a 10 wt% solution of MAO in toluene and stirred under 1 atmosphere ethylene.

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Ethylene uptake and formation of a polyethylene precipitate were observed. After 6.33 minutes, the mixture was quenched by the addition of acetone (50 mL), methanol (50 mL) and 6 N aqueous HCl (100 mL). The swollen polyethylene which separated was isolated by vacuum filtration, then dried at 80° C *in vacuo* for several hours. 860 mg of a white powdery polyethylene was isolated (87,000 TO/h). ^1H NMR: 15 branches/1000 carbon atoms. GPC: $M_n = 76,000$; $M_w/M_n = 2.6$

Example 73

10 Polymerization of ethylene using a catalyst generated *in situ* from tetrakis(2,6-dimethylphenyl)oxalamidine, bis(1,5-cyclooctadiene)nickel(0) and HB(Ar)₄ (Ar = 3,5-bis(trifluoromethyl)phenyl).

15 A 250 mL pear-shaped Schlenk flask equipped with a magnetic stir bar and capped with a septum was charged with 8.0 mg of bis(1,5-cyclooctadiene)nickel(0), 19 mg of N^1,N^2,N^3,N^4 -tetrakis(2,6-dimethylphenyl)oxalamidine, and 33 mg of the ether solvate of HB(Ar)₄. The flask was evacuated and refilled with ethylene, then charged with 75 mL of dry, deoxygenated toluene. The yellow solution which resulted was stirred under ethylene at 0° C for 30 minutes, then warmed to 25° C and 20 stirred for another 30 minutes under ethylene before being quenched by addition of acetone (50 mL), and methanol (50 mL). The polyethylene which separated was isolated by vacuum filtration and washed with water, methanol and acetone, then dried under reduced pressure (0.5 mm Hg) for 14 hours to give 0.70 g of an elastic white polyethylene (average 860 TO/h). 25 ^1H NMR: 83 branches/1000 carbon atoms. GPC: $M_n = 173,000$; $M_w/M_n = 2.8$.

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Example 74

Polymerization of ethylene with the nickel dibromide complex of 1,4-dimethyl-2,3-bis(2,6-dimethylphenylimino)piperazine in the presence of MAO.

5 A 250 mL pear-shaped Schlenk flask equipped with a magnetic stir bar and capped with a septum was charged with 10.4 mg of the nickel dibromide complex of 1,4-dimethyl-2,3-bis(2,6-dimethylphenylimino)piperazine. The flask was evacuated and refilled with ethylene, then charged with 75 mL of dry, deoxygenated toluene. The 10 resultant suspension was cooled to 0° C and allowed to equilibrate with 1 atmosphere ethylene for 15 minutes, then treated with 4.0 mL of a 10 wt% solution of MAO in toluene. The yellow solution which resulted was stirred under ethylene at 0° C for 1 hour, and then quenched by the addition of acetone (50 mL), methanol (50 mL) and 6 N aqueous HCl (100 mL). The 15 swollen polyethylene which separated was isolated by vacuum filtration and washed with water, methanol and acetone, then dried under reduced pressure (0.5 mm Hg) for 14 hours to give 1.3 g of a clear elastic polyethylene (2531 TO/h). ^1H NMR: 91 branches/1000 carbon atoms. GPC: $M_n = 127,000$; $M_w/M_n = 1.3$.

20

Example 75

Ethylene polymerization using complex XXXV.

25 A flame dried Schlenk flask equipped with a stir bar and a rubber septum was charged with 50 ml of methylene chloride and 50 mg of the palladium complex XXXV. The flask was placed under an ethylene atmosphere (1 atmosphere). The mixture was left to stir for 20 hours. Acetone and methanol were added to quench the reaction and precipitate the polymer. The polymer was collected and dried *in vacuo* resulting in 2.6 g of tacky polymer. NMR spectroscopic analysis is consistent with the 30 preparation of an ethylene homopolymer. ^1H NMR: highly branched

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polyethylene. GPC: $M_n = 34,000$, $M_w/M_n = 2.5$. DSC: $T_m = -39$ °C, $T_g = -69$ °C.

Example 76

5 Propylene polymerization using complex XXXV.

A flame dried Schlenk flask equipped with a stir bar and a rubber septum was charged with 50 ml of methylene chloride and 50 mg of the palladium complex XXXV. The flask was placed under a propylene atmosphere (1 atmosphere). The mixture was left to stir for 20 hours. 10 Acetone and methanol were added to quench the reaction and precipitate the polymer. The polymer was collected and dried *in vacuo* resulting in 580 mg of tacky polymer. 1H NMR: 192 branch points/1000 carbon atoms. GPC: $M_n = 17,000$, $M_w/M_n = 2.08$. DSC: $T_g = -53$ °C.

15 Example 77

Ethylene/vinyl ethylene carbonate copolymerization using complex XXXV.

A 200 mL flame dried pear-shaped Schlenk flask equipped with a magnetic stir bar and capped with a septum was charged with (2,6-diisopropylphenylimino)-[1,4]dithiane Pd(II) catalyst XXXV (100 mg) in an 20 argon filled glove box. Upon removal from the glove box, the flask was evacuated and backfilled with ethylene. The catalyst was dissolved in CH_2Cl_2 (25 mL) and immediately treated with vinyl ethylene carbonate (5mL). The resulting orange solution was stirred at 23 °C under an ethylene atmosphere (1 atm) for 20 hours. A small amount of polymer had 25 precipitated out of solution. The polymerization was quenched with MeOH and acetone leaving gray oil adhering to the walls of the flask. The polymer was washed several times with acetone and MeOH to remove any remaining monomer. The polymer was dissolved in CH_2Cl_2 and transferred to a storage jar. The solvent was left to evaporate and the resulting oily 30 polymer was dried *in vacuo* at ~80 °C for 3 days to afford a tacky solid (2.15

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g, 1100 TO). ^1H NMR was consistent with a copolymer containing approximately 96.5 weight % ethylene and 3.5 weight % vinyl ethylene carbonate monomer units.; M_n 40,200 g/mol ; M_w 92,100 g/mol ; DSC T_g -68 °C, T_m -38 °C.

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Example 78

Ethylene/vinyl ethylene carbonate copolymerization using complex XXXV.

A 200 mL flame dried pear-shaped Schlenk flask equipped with a magnetic stir bar and capped with a septum was charged with (2,6-di-isopropylphenylimino)-[1,4]dithiane Pd(II) catalyst XXXV (100 mg) in an argon filled glove box. Upon removal from the glove box, the flask was evacuated and backfilled with ethylene. The catalyst was dissolved in CH_2Cl_2 (20 mL) and immediately treated with vinyl ethylene carbonate (10mL). The resulting orange solution was stirred at 23 °C under an ethylene atmosphere (1 atm) for 28 hours. A small amount of polymer had precipitated out of solution. The polymerization was quenched with MeOH and acetone leaving gray oil adhering to the walls of the flask. The polymer was dissolved in CH_2Cl_2 and transferred to a storage jar. The solvent was left to evaporate and the resulting oily polymer was washed several times with acetone and MeOH to remove any remaining monomer and dried *in vacuo* at ~80 °C for 1 day to afford a tacky solid (1.15 g, 613 TO). ^1H NMR was consistent with a copolymer containing approximately 95.5 weight % ethylene and 4.5 weight % vinyl ethylene carbonate monomer units.; M_n 15,400 g/mol ; M_w 96,000 g/mol ; DSC T_g -64 °C, T_m -31 °C.

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Example 79

Preparation of the nickel dibromide complex of 2,3-bis(2,6-diisopropylphenylimino)-[1,4]dioxane.

A Schlenk flask equipped with a magnetic stir bar was charged 5 with 100 mg of 2,3-bis(2,6-diisopropylphenylimino)-[1,4]dioxane (0.25 mmol) and 71 mg of (1,2-dimethoxyethane)nickel(II) dibromide (0.23 mmol) under an argon atmosphere. Dry, deoxygenated dichloromethane (15 mL) was added and the mixture was stirred under an argon atmosphere, turning red-brown within about 10 minutes. After 2 hours, the red/orange solution 10 was transferred via filter cannula to a new flame dried Schlenk to remove trace amount of unreacted (1,2-dimethoxyethane)nickel(II) dibromide. The CH_2Cl_2 was removed *in vacuo*. The resulting red-brown solid was washed with 2x10 mL of hexane and the solid was dried *in vacuo* for several hours affording 80 mg of a brown solid.

Example 80

Preparation of 2,3-bis(2,6-dimethylphenylimino)-2,3-dihydroimidazo[2,1-*b*]thiazole.

A 50 mL round bottom flask equipped with a magnetic stir bar and a reflux condenser capped by a nitrogen inlet was charged with 752 mg of 20 N^1,N^2 -bis(2,6-dimethylphenyl)oxalodiimidoyl dichloride, 200 mg of sodium hydride (60% mineral oil dispersion), 5.0 mL of dry tetrahydrofuran, and 250 mg of 2-mercaptoimidazole. The mixture was heated at reflux for 120 minutes. After cooling, the mixture was diluted with water and dichloromethane, and the organic layer was separated and concentrated to afford a yellow-orange oil. Column chromatography (SiO_2 , Merck Grade 25 9385 230-400 mesh, 60 Å; 12 v% ethyl acetate in hexane) afforded 487 mg of a yellow-orange solid. Recrystallization from heptane gave 366 mg yellow-orange prisms. Field desorption mass spectrometry showed a parent ion peak at 360 m/z.

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Example 81

Preparation of N^1, N^2 -bis(2,6-dimethylphenyl)ethanediimidatoselenoic acid diphenyl ester.

A 50 mL round bottom flask equipped with a magnetic stir bar and a reflux condenser capped by a nitrogen inlet was charged with 961 mg of N^1, N^2 -bis(2,6-dimethylphenyl)oxalodiimoyl dichloride, 287 mg of sodium hydride (60% mineral oil dispersion), 8.2 mL of dry tetrahydrofuran, and .068 mL of benzeneselenol. The mixture was heated at reflux for 45 minutes. After cooling, the mixture was diluted with water and diethyl ether. The ether layer was separated and washed again with water, then concentrated *in vacuo* to afford a yellow-orange crystalline solid. The solid was dissolved in hot hexane, then filtered, and then reconcentrated. Recrystallization from heptane afforded 745 mg orange prisms, 1st crop. Field desorption mass spectrometry showed a parent ion cluster of peaks from 570-578 m/z. 1H NMR (300 MHz, $CDCl_3$, chemical shifts in ppm relative to TMS at 0 ppm): 1.95 (12 p, s), 6.75 (6p, app s), 7.02-7.20 (6p, m), 7.39-7.48 (4p, m).

Example 82

20 Polymerization of ethylene using a catalyst generated *in situ* from N^1, N^2 -bis(2,6-dimethylphenyl)ethanediimidatoselenoic acid diphenyl ester, bis(1,5-cyclooctadiene)nickel(0) and $HB(Ar)_4$ ($Ar = 3,5$ -bis(trifluoromethyl)phenyl).

A 250 mL pear-shaped Schlenk flask equipped with a magnetic stir bar and capped with a septum was charged with 5 mg of bis(1,5-cyclooctadiene)nickel(0), 10 mg of N^1, N^2 -bis(2,6-dimethylphenyl)ethanediimidatoselenoic acid diphenyl ester, and 25 mg of the ether solvate of $HB(Ar)_4$. The flask was evacuated and refilled with ethylene, then charged with 45 mL of dry, deoxygenated toluene. The yellow solution which resulted was stirred under ethylene at 21° C for 10 minutes, then quenched by addition of methanol (50 mL). The

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polyethylene which separated was isolated by vacuum filtration and washed with methanol, then dried under reduced pressure (0.5 mm Hg) for 14 hours to give 0.060 g of an elastic blue-green polyethylene. ^1H NMR: 24 branches/1000 carbon atoms. GPC: $M_n = 181,000$; $M_w/M_n = 3.5$.

5

Example 83

Preparation of the nickel dibromide complex of 2,3-bis(2,6-dimethylphenylimino)-2,3-dihydroimidazo[2,1-*b*]thiazole.

A 50 mL Schlenk flask equipped with a magnetic stir bar and capped with a septum was charged with 141 mg of 2,3-bis(2,6-dimethylphenylimino)-2,3-dihydroimidazo[2,1-*b*]thiazole and 110 mg of (1,2-dimethoxyethane)nickel(II) dibromide under an inert atmosphere. Dry, deoxygenated dichloromethane (5 mL) was added and the mixture was stirred under an argon atmosphere. After 1 hour, another 5 mL of dichloromethane was added. The mixture was stirred another 16 hours at 21° C, then diluted with 10 mL of dry, deoxygenated hexane and stirred another 3 hours. The supernatant was removed via a filter paper-tipped cannula, and the residue dried *in vacuo* at 1 mm Hg to afford 66 mg of a brown microcrystalline solid.

20

Example 84

Polymerization of ethylene with the nickel dibromide complex 2,3-bis(2,6-dimethylphenylimino)-2,3-dihydroimidazo[2,1-*b*]thiazole in the presence of MAO.

A 200 mL pear-shaped Schlenk flask equipped with a magnetic stir bar and capped with a septum was charged with 2.5 mg of the nickel dibromide complex of 2,3-bis(2,6-dimethylphenylimino)-2,3-dihydroimidazo[2,1-*b*]thiazole. The flask was evacuated and refilled with ethylene, then charged with 75 mL of dry, deoxygenated toluene. The resultant suspension allowed to equilibrate with 1 atmosphere ethylene at

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21 °C for 15 minutes, then treated with 200 μ L of a 10 wt% solution of MAO in toluene and stirred under 1 atmosphere ethylene. After 21 min, the reaction was quenched by the addition of acetone (50 mL), methanol (50 mL) and 6 N aqueous HCl (100 mL). The swollen polyethylene which separated was isolated by vacuum filtration and washed with water, methanol and acetone, then dried under reduced pressure (0.05-0.1 mm Hg) for 24 hours to give 198 mg of a white polyethylene. 1 H NMR: 13 branches/1000 carbon atoms. GPC: bimodal, with M_n = 23,000; M_p = 366,000; M_w/M_n = 13.5.

10

Example 85

Preparation of the nickel dibromide complex of N^1,N^2,N^3,N^4 -tetrakis(2,6-dimethylphenyl)oxalamidine.

A 50 mL Schlenk flask equipped with a magnetic stir bar and capped with a septum was charged with 100 mg of N^1,N^2,N^3,N^4 -tetrakis(2,6-dimethylphenyl)oxalamidine and 55 mg of (1,2-dimethoxyethane)nickel(II) dibromide under an inert atmosphere. Dry, deoxygenated dichloromethane (5 mL) was added and the mixture was stirred under an argon atmosphere. After 1 hour, another 5 mL of dichloromethane was added. The mixture was stirred another 16 hours at 21° C, then diluted with 10 mL of dry, deoxygenated hexane and stirred another 3 hours. The supernatant was removed via a filter paper-tipped cannula, and the residue dried *in vacuo* at 1 mm Hg to afford 95 mg of light green crystals.

25

Example 86

Polymerization of ethylene with the nickel dibromide complex of N^1,N^2,N^3,N^4 -tetrakis(2,6-dimethylphenyl)oxalamidine in the presence of MAO.

A 200 mL pear-shaped Schlenk flask equipped with a magnetic stir bar and capped with a septum was charged with 2.4 mg of nickel

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dibromide complex of N^1,N^2,N^3,N^4 -tetrakis(2,6-dimethylphenyl)oxalamidine. The flask was evacuated and refilled with ethylene, then charged with 75 mL of dry, deoxygenated toluene. The resultant suspension allowed to equilibrate with 1 atmosphere ethylene at 21 °C for 15 minutes, then treated 5 with 4.0 mL of a 10 wt% solution of MAO in toluene and stirred under 1 atmosphere ethylene. After 30 min, the reaction was quenched by the addition of acetone (50 mL), methanol (50 mL) and 6 N aqueous HCl (100 mL). The swollen polyethylene which separated was isolated by vacuum filtration and washed with water, methanol and acetone, then dried under 10 reduced pressure (0.05-0.1 mm Hg) for 24 hours to give 743 mg of a white polyethylene. ^1H NMR: 112 branches/1000 carbon atoms. GPC: $M_n = 330,000$; $M_w/M_n = 1.4$.

Example 87

15 Copolymerization of ethylene and 1-pentene with the nickel dibromide complex of 2,3-bis(2,6-dimethylphenylimino)-[1,4]dithiane in the presence of 20 MAO.

A 200 mL pear-shaped Schlenk flask equipped with a magnetic stir bar and capped with a septum was charged with 0.5 mL of a stock solution 20 of 12.4 mg of the nickel dibromide complex of 2,3-bis(2,6-dimethylphenylimino)-[1,4]dithiane in 10.0 mL dichloromethane. The flask was evacuated and refilled with ethylene, then charged with 100 mL of dry, deoxygenated toluene, and 5.0 mL 1-pentene. The resultant suspension was cooled to 0 °C and allowed to equilibrate with 1 atmosphere ethylene 25 for 15 minutes, then treated with 4.0 mL of a 10 wt% solution of MAO in toluene and stirred under 1 atmosphere ethylene. After 45 minutes, the mixture was quenched by the addition of acetone (50 mL), methanol (50 mL) and 6 N aqueous HCl (100 mL). The swollen copolymer which separated was isolated by vacuum filtration and washed with water, 30 methanol and acetone, then dried under reduced pressure (255 mm Hg) at

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100 °C for 24 hours to obtain 2.0 g of white copolymer. ^1H NMR: 24 branches/1000 carbon atoms. ^{13}C NMR: 7.6 methyl branches/1000 carbons, 1.2 ethyl branches/1000 carbons, 9.1 propyl branches/1000 carbons, 2.1 butyl branches/1000 carbons, 3.4 pentyl and higher alkyl branches/1000 carbons. GPC: $M_n = 274,000$; $M_w/M_n = 2.3$.

Example 88

Copolymerization of ethylene and 1-heptene with the nickel dibromide complex of 2,3-bis(2,6-dimethylphenylimino)-[1,4]dithiane in the presence of MAO.

A 200 mL pear-shaped Schlenk flask equipped with a magnetic stir bar and capped with a septum was charged with 0.5 mL of a stock solution of 12.4 mg of the nickel dibromide complex of 2,3-bis(2,6-dimethylphenylimino)-[1,4]dithiane in 10.0 mL dichloromethane. The flask was evacuated and refilled with ethylene, then charged with 100 mL of dry, deoxygenated toluene, and 5.0 mL 1-heptene. The resultant suspension was cooled to 0 °C and allowed to equilibrate with 1 atmosphere ethylene for 15 minutes, then treated with 4.0 mL of a 10 wt% solution of MAO in toluene and stirred under 1 atmosphere ethylene. After 33 minutes, the mixture was quenched by the addition of acetone (50 mL), methanol (50 mL) and 6 N aqueous HCl (100 mL). The swollen copolymer which separated was isolated by vacuum filtration and washed with water, methanol and acetone, then dried under reduced pressure (255 mm Hg) at 100 °C for 24 hours to obtain 1.25 g of white copolymer. ^1H NMR: 19 branches/1000 carbon atoms. ^{13}C NMR: 5.9 methyl branches/1000 carbons, less than 1 ethyl branch/1000 carbons, less than 1 propyl branch/1000 carbons, 1.8 butyl branches/1000 carbons, 11.5 pentyl and higher alkyl branches/1000 carbons. GPC: $M_n = 223,000$; $M_w/M_n = 2.3$.

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Example 89

Polymerization of 1-hexene with the nickel dibromide complex of 2,3-bis(2,6-dimethylphenylimino)-[1,4]dithiane in the presence of MAO.

A 22 mL vial equipped with a magnetic stir bar and capped by a 5 septum was sequentially charged with 1.8 mg of the nickel dibromide complex of 2,3-bis(2,6-dimethylphenylimino)-[1,4]dithiane, 4.0 mL 1-hexene, and 2.0 mL of a 10 wt% solution of MAO in toluene, under Ar. The resultant violet mixture thickened noticeably within minutes. After 34 min, the reaction was quenched with acetone, methanol and 6 N aq HCl, and the 10 polyhexene which separated was filtered off and dried in vacuo (0.4 mm Hg) to obtain 428 mg of an elastic polyhexene. ^1H NMR: 173 branches/1000 carbon atoms. GPC: $M_n = 92,000$; $M_w/M_n = 2.0$.

15

Example 90

Polymerization of 1-hexene with the nickel dibromide complex 2,3-bis(2,6-dimethylphenylimino)-2,3-dihydroimidazo[2,1-*b*]thiazole in the presence of MAO.

A 22 mL vial equipped with a magnetic stir bar and capped by a 20 septum was sequentially charged with 2.1 mg of the nickel dibromide complex 2,3-bis(2,6-dimethylphenylimino)-2,3-dihydroimidazo[2,1-*b*]thiazole, 4.0 mL 1-hexene, and 2.0 mL of a 10 wt% solution of MAO in toluene, under Ar. The resultant dark purple-brown mixture thickened noticeably within 10-20 minutes. After 53 min, the reaction was quenched 25 with acetone, methanol and 6 N aq HCl, and the polyhexene which separated was filtered off and dried in vacuo (0.4 mm Hg) to obtain 283 mg of an elastic polyhexene. ^1H NMR: 110 branches/1000 carbon atoms. GPC: $M_n = 91,000$; $M_w/M_n = 1.9$.

30

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Example 91

Polymerization of 1-hexene with the nickel dibromide complex of 1,4-dimethyl-2,3-bis(2,6-dimethylphenylimino)piperazine in the presence of MAO.

5 A 22 mL vial equipped with a magnetic stir bar and capped by a septum was sequentially charged with 2.1 mg of nickel dibromide complex of 1,4-dimethyl-2,3-bis(2,6-dimethylphenylimino)piperazine, 4.0 mL 1-hexene, and 2.0 mL of a 10 wt% solution of MAO in toluene, under Ar. The resultant clear yellow solution was stirred at 23 °C for 400 min, then the 10 reaction was quenched with acetone, methanol and 6 N aq HCl, and the polyhexene which separated was filtered off and dried in vacuo (0.4 mm Hg) to obtain 408 mg of an elastic polyhexene. ^1H NMR: 90 branches/1000 carbon atoms. GPC: $M_n = 47,000$; $M_w/M_n = 1.7$.

15

Obviously, numerous modifications and variations of the present invention are possible in light of the above teachings. It is therefore to be 20 understood that within the scope of the appended Claims, the invention may be practiced otherwise than as specifically described herein.

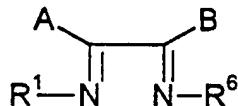
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CLAIMS

We claim:

5 1. A batch or continuous process for the polymerization of olefins, comprising contacting one or more monomers selected from compounds of the formula $RCH=CHR^8$ with a catalyst comprising (a) a suitable metal precursor and (b) a ligand of the formula X, and optionally (c) a Bronsted or Lewis acid,

10



X

wherein R and R⁸ each, independently, represent hydrogen, hydrocarbyl, or fluoroalkyl, and may be linked to form a cyclic olefin; R¹ and R⁶ are each, independently, hydrocarbyl, substituted hydrocarbyl, or silyl; N represents nitrogen;

15 A and B are each, independently, a heteroatom connected monoradical wherein the connected heteroatom is selected from Group 15 or 16; in addition, A and B may be linked by a bridging group.

20 2. The process of Claim 1 wherein the suitable metal precursor is a Ni compound.

25 3. The process of Claim 1 wherein the suitable metal precursor is a Co compound.

4. The process of Claim 1 wherein the suitable metal precursor is a Pd compound.

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5. The process of Claim 1 wherein the suitable metal precursor is an Fe compound.

5 6. The process of Claim 1 wherein the monomer of the formula RCH=CHR⁸ is selected from ethylene, propylene, 1-butene, 1-hexene, and 1-octene.

10 7. The process of Claim 2 wherein the monomer of the formula RCH=CHR⁸ is selected from ethylene, propylene, 1-butene, 1-hexene, and 1-octene.

15 8. The process of Claim 4 wherein the monomers further comprise methyl acrylate.

9. The process of Claim 4 wherein the monomers further comprise vinyl acetate.

20 10. The process of Claim 2 wherein a Lewis acid is utilized and said Lewis acid is diethylaluminum chloride.

11. The process of Claim 2 wherein a Lewis acid is utilized and said Lewis acid is methylalumoxane.

25 12. The process of Claim 2 wherein R¹ and R⁶ are 2,6-dimethylphenyl.

13. The process of Claim 2 wherein R¹ and R⁶ are 2,6-diisopropylphenyl.

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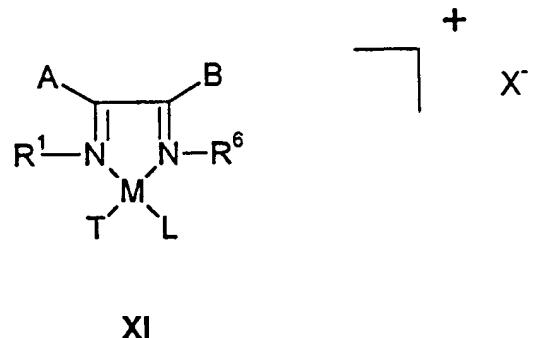
14. The process of Claim 2, wherein the polymerization is conducted in an inert solvent.

15. The process of Claim 14, wherein the inert solvent is selected
5 from linear or branched hydrocarbons, toluene, or mixtures thereof.

16. The process of Claim 2, wherein the catalyst is attached to a solid support.

10 17. The process of Claim 2 wherein the temperature is between about 25 °C and about 150 °C, and the pressure is between about 1 atm and about 1000 atm.

15 18. A batch or continuous process for the polymerization of olefins, comprising contacting one or more monomers of the formula RCH=CHR⁸ with a catalyst of formula XI,



20 wherein R and R⁸ each, independently, represent hydrogen, hydrocarbyl, or fluoroalkyl, and may be linked to form a cyclic olefin; R¹ and R⁶ each, independently, represent hydrocarbyl, substituted hydrocarbyl, or silyl;

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A and B are each, independently, a heteroatom connected mono-radical wherein the connected heteroatom is selected from Group 15 or 16; in addition, A and B may be linked by a bridging group;

T represents a hydrogen or a hydrocarbyl;

5 L represents a mono-olefin or a neutral Lewis base wherein the coordinated atom is nitrogen, oxygen, or sulfur;

M represents Ni(II), Pd(II), Co(II), or Fe(II);

N represents nitrogen; and

X⁻ is a weakly coordinating anion.

10

19. The process of Claim 18 wherein M is Ni(II).

20. The process of Claim 18 wherein M is Co(II).

15

21. The process of Claim 18 wherein M is Pd(II).

22. The process of Claim 18 wherein M is Fe(II).

20

23. The process of Claim 18 wherein the monomer of the formula RCH=CHR⁸ is selected from ethylene, propylene, 1-butene, 1-hexene, and 1-octene.

25

24. The process of Claim 19 wherein the monomer of the formula RCH=CHR⁸ is selected from ethylene, propylene, 1-butene, 1-hexene, and 1-octene.

25. The process of Claim 21 wherein the monomers further comprise methyl acrylate.

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26. The process of Claim 21 wherein the monomers further comprise vinyl acetate.

27. The process of Claim 19 wherein R¹ and R⁶ are 2,6-
5 dimethylphenyl.

28. The process of Claim 19 wherein R¹ and R⁶ are 2,6-
diisopropylphenyl.

10 29. The process of Claim 19, wherein the polymerization is conducted in an inert solvent.

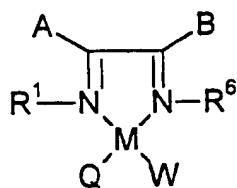
30. The process of Claim 29, wherein the inert solvent is selected from linear or branched hydrocarbons, toluene, or mixtures thereof.

15 31. The process of Claim 19, wherein the catalyst is attached to a solid support.

20 32. The process of Claim 19 wherein the temperature is between about 25 °C and about 150 °C, and the pressure is between about 1 atm and about 1000 atm.

25 33. A batch or continuous process for the polymerization of olefins, comprising contacting one or more monomers of the formula RCH=CHR⁸ with a catalyst formed by combining a compound of formula XII:

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XII

with a compound Y,

wherein R and R⁸ each, independently, represent a hydrogen, a hydrocarbyl, or a fluoroalkyl, and may be linked to form a cyclic olefin;

5 R¹ and R⁶ each, independently, represent hydrocarbyl, substituted hydrocarbyl, or silyl;

A and B are each, independently, a heteroatom connected mono-radical wherein the connected heteroatom is selected from Group 15 or 16; in addition, A and B may be linked by a bridging group;

10 Q represents an alkyl, chloride, iodide or bromide;

W represents an alkyl, chloride, iodide or bromide;

N represents nitrogen; and

M represents Ni(II), Pd(II), Co(II), or Fe(II);

15 and Y is selected from the group consisting of a neutral Lewis acid capable of abstracting Q⁻ or W⁻ to form a weakly coordinating anion, a cationic Lewis acid whose counterion is a weakly coordinating anion, and a Bronsted acid whose conjugate base is a weakly coordinating anion.

34. The process of Claim 33 wherein M is Ni(II).

20

35. The process of Claim 33 wherein M is Co(II).

36. The process of Claim 33 wherein M is Pd(II).

25

37. The process of Claim 33 wherein M is Fe(II).

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38. The process of Claim 33 wherein the monomer of the formula RCH=CHR⁸ is selected from ethylene, propylene, 1-butene, 1-hexene, and 1-octene.

5 39. The process of Claim 34 wherein the monomer of the formula RCH=CHR⁸ is selected from ethylene, propylene, 1-butene, 1-hexene, and 1-octene.

10 40. The process of Claim 34 wherein Y is a Lewis acid and said Lewis acid is diethylaluminum chloride.

41. The process of Claim 34 wherein Y is a Lewis acid and said Lewis acid is methylalumoxane.

15 42. The process of Claim 34 wherein R¹ and R⁶ are 2,6-dimethylphenyl.

43. The process of Claim 34 wherein R¹ and R⁶ are 2,6-diisopropylphenyl.

20 44. The process of Claim 34, wherein the polymerization is conducted in an inert solvent.

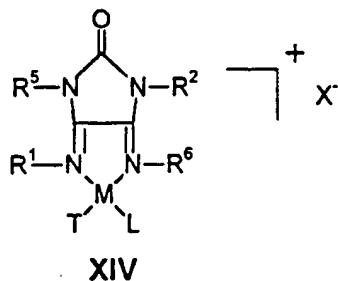
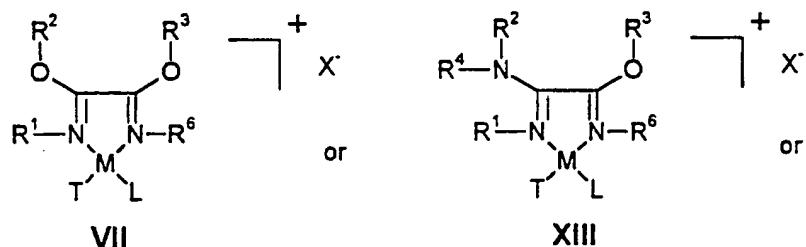
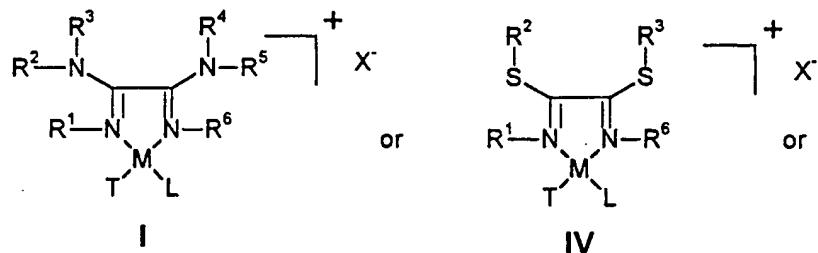
25 45. The process of Claim 44, wherein the inert solvent is selected from linear or branched hydrocarbons, toluene, or mixtures thereof.

46. The process of Claim 34, wherein the catalyst is attached to a solid support.

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47. The process of Claim 34 wherein the temperature is between about 25 °C and about 150 °C, and the pressure is between about 1 atm and about 1000 atm.

5 48. The process of Claim 18 wherein the catalyst of formula XI is selected from:



10 wherein R¹ and R⁶ each, independently, represent hydrocarbyl, substituted hydrocarbyl, or silyl;

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R^2 , R^3 , R^4 and R^5 each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl; in addition, any two of R^2 , R^3 , R^4 , and R^5 may collectively form a bridging group, provided that when the catalyst is of formula IV or VII, the bridging group is not a substituted sulfur atom or a substituted phosphorous atom;

5 T represents a hydrogen or hydrocarbyl;

L represents a mono-olefin or a neutral Lewis base wherein the coordinated atom is nitrogen, oxygen, or sulfur;

M represents Ni(II), Pd(II), Co(II) or Fe(II);

10 N represents nitrogen; O represents oxygen; S represents sulfur;

 and

X^- is a weakly coordinating anion.

49. The process of Claim 48, wherein the catalyst is of the
15 formula I.

50. The process of Claim 48, wherein the catalyst is of the
 formula IV.

20 51. The process of Claim 48, wherein the catalyst is of the
 formula VII.

52. The process of Claim 48, wherein the catalyst is of the
 formula XIII.

25 53. The process of Claim 48, wherein the catalyst is of the
 formula XIV.

54. The process of Claim 49 wherein M is Ni(II).

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55. The process of Claim 50 wherein M is Ni(II).
56. The process of Claim 51 wherein M is Ni(II).
- 5 57. The process of Claim 52 wherein M is Ni(II).
58. The process of Claim 53 wherein M is Ni(II).
59. The process of Claim 48 wherein M is Ni(II).
- 10 60. The process of Claim 48 wherein M is Co(II).
61. The process of Claim 48 wherein M is Pd(II).
- 15 62. The process of Claim 48 wherein M is Fe(II).
63. The process of Claim 59 wherein the monomer of the formula RCH=CHR⁸ is selected from ethylene, propylene, 1-butene, 1-hexene, and 1-octene.
- 20 64. The process of Claim 59 wherein R¹ and R⁶ are 2,6-dimethylphenyl.
65. The process of Claim 59 wherein R¹ and R⁶ are 2,6-diisopropylphenyl.
- 25 66. The process of Claim 59, wherein the polymerization is conducted in an inert solvent.

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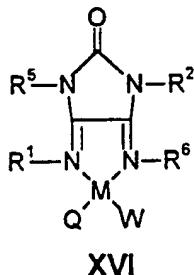
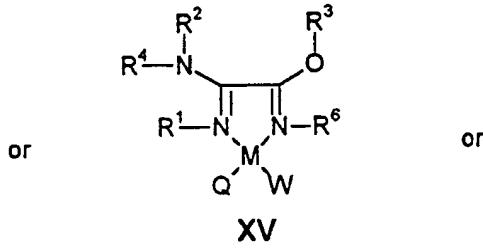
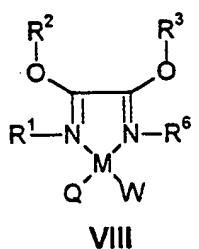
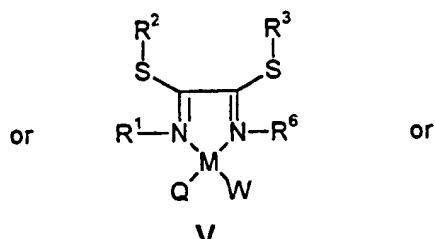
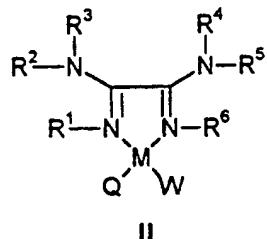
67. The process of Claim 66, wherein the inert solvent is selected from linear or branched hydrocarbons, toluene, or mixtures thereof.

68. The process of Claim 59, wherein the catalyst is attached to a
5 solid support.

69. The process of Claim 59 wherein the temperature is between about 25 °C and about 150 °C, and the pressure is between about 1 atm and about 1000 atm.

10 70. The process of Claim 33 wherein the catalyst of formula XII is selected from:

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wherein R and R⁸ each, independently, represent hydrogen,

hydrocarbyl, or fluoroalkyl, and may be linked to form a cyclic olefin;

R¹ and R⁶ each, independently, represent hydrocarbyl, substituted

hydrocarbyl, or silyl;

R², R³, R⁴ and R⁵ each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl; in addition, any two of R², R³, R⁴, and R⁵ may collectively form a bridging group, provided that when the catalyst is of formula V or VIII, the bridging group is not a substituted sulfur atom or a substituted phosphorous atom;

Q represents a hydrocarbyl, chloride, iodide or bromide;

W represents a hydrocarbyl, chloride, iodide or bromide;

M represents Ni(II), Pd(II), Co(II), or Fe(II); and

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N represents nitrogen.

71. The process of Claim 70, wherein the catalyst is of the formula II.

5

72. The process of Claim 70, wherein the catalyst is of the formula V.

10 73. The process of Claim 70, wherein the catalyst is of the formula VIII.

74. The process of Claim 70, wherein the catalyst is of the formula XV.

15 75. The process of Claim 70, wherein the catalyst is of the formula XVI.

76. The process of Claim 71 wherein M is Ni(II).

20 77. The process of Claim 72 wherein M is Ni(II).

78. The process of Claim 73 wherein M is Ni(II).

79. The process of Claim 74 wherein M is Ni(II).

25

80. The process of Claim 75 wherein M is Ni(II).

81. The process of Claim 70 wherein M is Ni(II).

30

82. The process of Claim 70 wherein M is Co(II).

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83. The process of Claim 70 wherein M is Pd(II).

84. The process of Claim 70 wherein M is Fe(II).

5

85. The process of Claim 81 wherein the monomer of the formula $RCH=CHR^8$ is selected from ethylene, propylene, 1-butene, 1-hexene, and 1-octene.

10 86. The process of Claim 83 wherein the monomers further comprise methyl acrylate.

87. The process of Claim 83 wherein the monomers further comprise vinyl acetate.

15

88. The process of Claim 81 wherein R^1 and R^6 are 2,6-dimethylphenyl.

20 89. The process of Claim 81 wherein R^1 and R^6 are 2,6-diisopropylphenyl.

90. The process of Claim 81, wherein the polymerization is conducted in an inert solvent.

25

91. The process of Claim 90, wherein the inert solvent is selected from linear or branched hydrocarbons, toluene, or mixtures thereof.

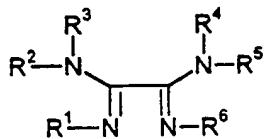
92. The process of Claim 81, wherein the catalyst is attached to a solid support.

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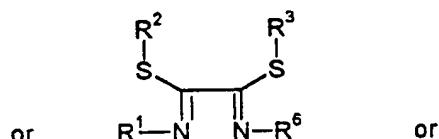
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93. The process of Claim 81, wherein the temperature is between about 25 °C and about 150 °C, and the pressure is between about 1 atm and about 1000 atm.

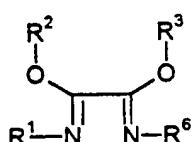
5 94. The process of Claim 1 wherein the suitable metal precursor is a nickel(0) compound and the ligand of formula X is selected from:



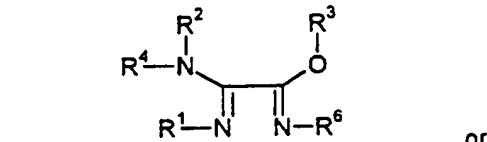
III



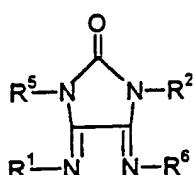
VI



IX



XVII



XVIII

and is combined with a Bronsted acid whose conjugate base is a weakly coordinating anion,

10 wherein R¹ and R⁶ each, independently, represent hydrocarbyl, substituted hydrocarbyl, or silyl;

R², R³, R⁴ and R⁵ each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl; in addition, any two of R², R³,

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R^4 , and R^5 may collectively form a bridging group, provided that when the ligand is of formula VI or IX, the bridging group is not a substituted sulfur atom or a substituted phosphorous atom.

5 95. The process of Claim 94 wherein the monomer of the formula $RCH=CHR^8$ is selected from ethylene, propylene, 1-butene, 1-hexene, and 1-octene.

10 96. The process of Claim 94 wherein R^1 and R^6 are 2,6-dimethylphenyl.

97. The process of Claim 94 wherein R^1 and R^6 are 2,6-diisopropylphenyl.

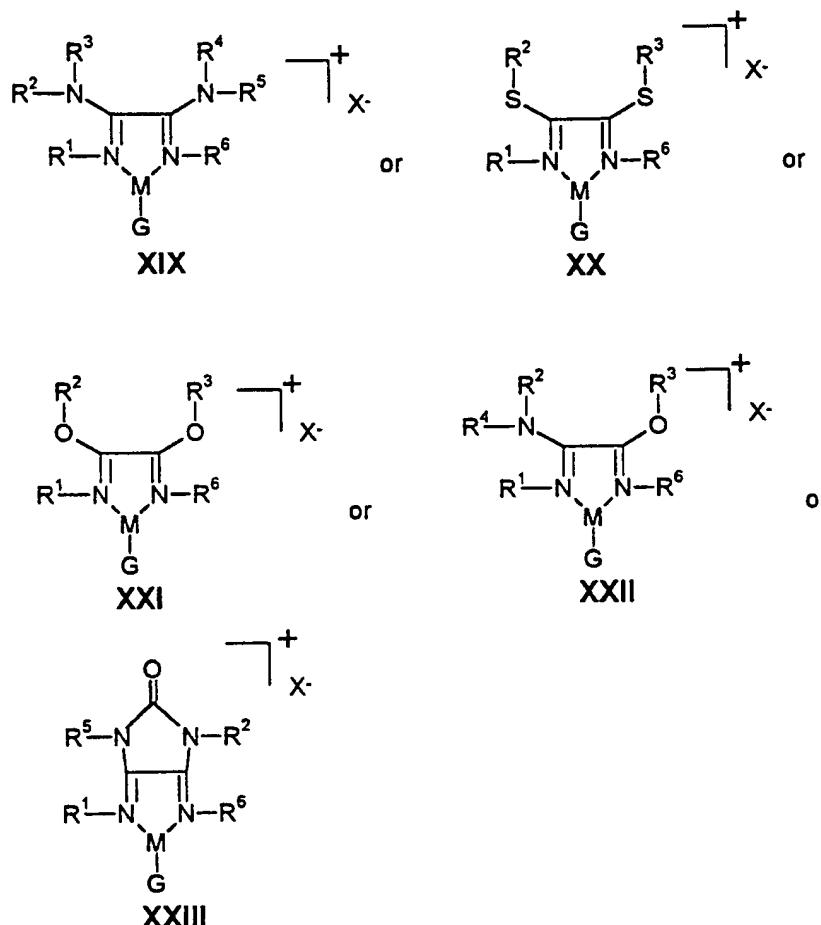
15 98. The process of Claim 94, wherein the polymerization is conducted in an inert solvent.

99. The process of Claim 98, wherein the inert solvent is selected from linear or branched hydrocarbons, toluene, or mixtures thereof.

20 100. The process of Claim 94, wherein the temperature is between about 25 °C and about 150 °C, and the pressure is between about 1 atm and about 1000 atm.

25 101. A batch or continuous process for the polymerization of olefins, comprising contacting one or more monomers of the formula $RCH=CHR^8$ with a catalyst of the formula:

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wherein R and R^8 each, independently, represent hydrogen, hydrocarbyl, or fluoroalkyl, and may be linked to form a cyclic olefin;

R^1 and R^6 each, independently, represent hydrocarbyl, substituted hydrocarbyl, or silyl;

5 R^2 , R^3 , R^4 and R^5 each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl; in addition, any two of R^2 , R^3 , R^4 , and R^5 may collectively form a bridging group, provided that when the catalyst is of formula XX or XXI, the bridging group is not a substituted sulfur atom or a substituted phosphorous atom;

10 G is a π -allyl or π -benzyl group;

M represents Ni(II), or Pd(II);

- 129 -

N represents nitrogen; and
X⁻ is a weakly coordinating anion.

102. The process of Claim 101, wherein the catalyst is of the
5 formula XIX.

103. The process of Claim 101, wherein the catalyst is of the
formula XX.

104. The process of Claim 101, wherein the catalyst is of the
formula XXI.

105. The process of Claim 101, wherein the catalyst is of the
formula XXII.

15 106. The process of Claim 101, wherein the catalyst is of the
formula XXIII.

107. The process of Claim 101 wherein M is Ni(II).

20 108. The process of Claim 102 wherein M is Ni(II).

109. The process of Claim 103 wherein M is Ni(II).

25 110. The process of Claim 104 wherein M is Ni(II).

111. The process of Claim 105 wherein M is Ni(II).

112. The process of Claim 106 wherein M is Ni(II).

30

- 130 -

113. The process of Claim 101 wherein X⁻ is BF₄⁻.

114. The process of Claim 101 wherein M is Pd(II).

5 115. The process of Claim 101 wherein X⁻ is PF₆⁻.

116. The process of Claim 107 wherein the monomer of the formula RCH=CHR⁸ is selected from ethylene, propylene, 1-butene, 1-hexene, and 1-octene.

10

117. The process of Claim 114 wherein the monomers further comprise methyl acrylate.

15

118. The process of Claim 114 wherein the monomers further comprise vinyl acetate.

119. The process of Claim 107 wherein R¹ and R⁶ are 2,6-dimethylphenyl.

20

120. The process of Claim 107 wherein R¹ and R⁶ are 2,6-diisopropylphenyl.

121. The process of Claim 107, wherein the polymerization is conducted in an inert solvent.

25

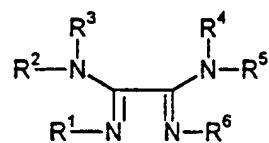
122. The process of Claim 121, wherein the inert solvent is selected from linear or branched hydrocarbons, toluene, or mixtures thereof.

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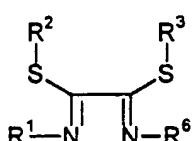
123. The process of Claim 107, wherein the catalyst is attached to a solid support.

124. The process of Claim 107 wherein the temperature is between 5 about 25 °C and about 150 °C, and the pressure is between about 1 atm and about 1000 atm.

125. A process for the production of linear α -olefins, comprising contacting ethylene with (a) a suitable nickel precursor, (b) a compound of 10 the formula:



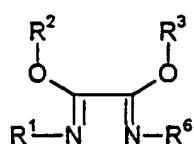
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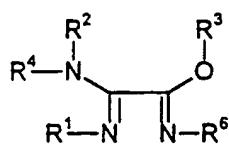
or

III

VI



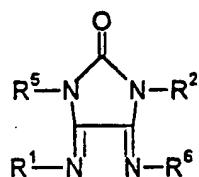
or



or

IX

XVII



XVIII

and, optionally (c) a Bronsted or Lewis acid, wherein

- 132 -

R^1 and R^6 each, independently, represent hydrocarbyl, substituted hydrocarbyl, or silyl;

R^2 , R^3 , R^4 and R^5 each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl; in addition, any two of R^2 , R^3 , 5 R^4 , and R^5 may collectively form a bridging group, provided that when the compound is of formula VI or IX, the bridging group is not a substituted sulfur atom or a substituted phosphorous atom;

N represents nitrogen; O represents oxygen; S represents sulfur.

10 126. The process of Claim 125, wherein the compound is of the formula III.

127. The process of Claim 125, wherein the compound is of the formula VI.

15 128. The process of Claim 125, wherein the compound is of the formula IX.

20 129. The process of Claim 125, wherein the compound is of the formula XVII.

130. The process of Claim 125, wherein the compound is of the formula XVIII.

25 131. The process of Claim 125 wherein R^1 and R^6 are phenyl.

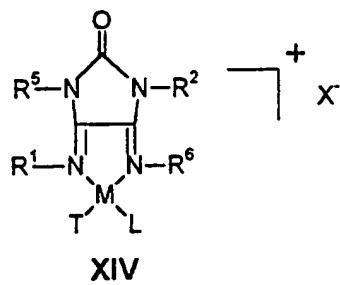
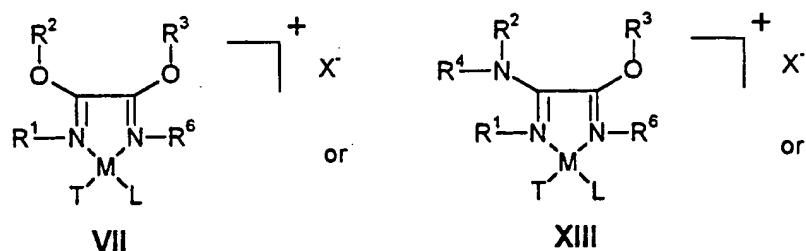
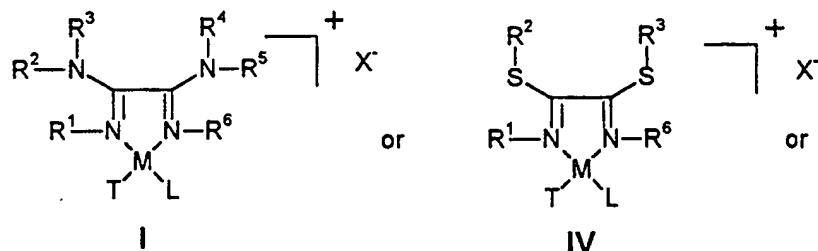
132. The process of Claim 125, wherein the process is conducted in an inert solvent.

- 133 -

133. The process of Claim 132, wherein the inert solvent is selected from linear or branched hydrocarbons, toluene, or mixtures thereof.

5 134. The process of Claim 125, wherein the temperature is
between about 25 °C and about 150 °C, and the pressure is between about
1 atm and about 1000 atm.

135. A process for the production linear α -olefins, comprising
10 contacting ethylene with a compound of the formula:



- 134 -

R^1 and R^6 each, independently, represent hydrocarbyl, substituted hydrocarbyl, or silyl;

R^2 , R^3 , R^4 and R^5 each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl; in addition, any two of R^2 , R^3 , 5 R^4 , and R^5 may collectively form a bridging group, provided that when the compound is of formula IV or VII, the bridging group is not a substituted sulfur atom or a substituted phosphorous atom;

T represents a hydrogen or hydrocarbyl;

10 L represents a mono-olefin or a neutral Lewis base wherein the coordinated atom is nitrogen, oxygen, or sulfur;

M represents $Ni(II)$;

N represents nitrogen; O represents oxygen; S represents sulfur;

and

X^- is a weakly coordinating anion.

15

136. The process of Claim 135, wherein the compound is of the formula I.

20

137. The process of Claim 135, wherein the compound is of the formula IV.

138. The process of Claim 135, wherein the compound is of the formula VII.

25

139. The process of Claim 135, wherein the compound is of the formula XIII.

140. The process of Claim 135, wherein the compound is of the formula XIV.

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- 135 -

141. The process of Claim 135, wherein R¹ and R⁶ are phenyl.

142. The process of Claim 135, wherein the process is conducted in an inert solvent.

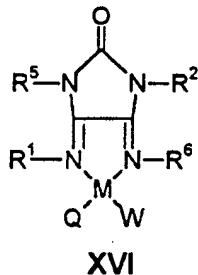
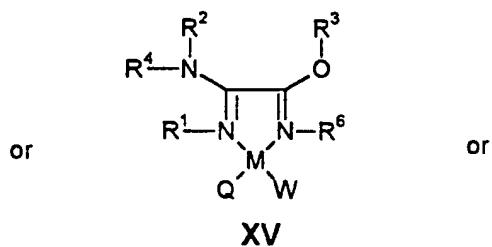
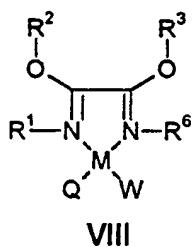
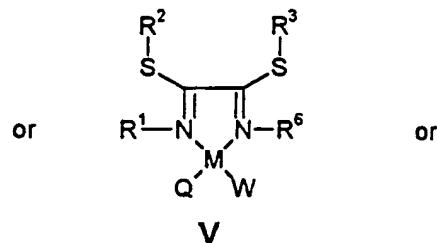
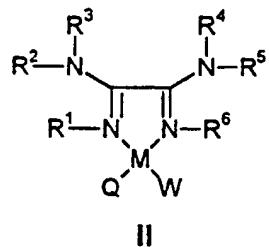
5

143. The process of Claim 142, wherein the inert solvent is selected from linear or branched hydrocarbons, toluene, or mixtures thereof.

10 144. The process of Claim 135 wherein the temperature is between about 25 °C and about 150 °C, and the pressure is between about 1 atm and about 1000 atm.

15 145. A process for the production of α -olefins, comprising contacting ethylene with a catalyst formed by combining a complex of the formula:

- 136 -



with a compound Y,

wherein R¹ and R⁶ each, independently, represent hydrocarbyl, substituted hydrocarbyl, or silyl;

5 R², R³, R⁴ and R⁵ each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl; in addition, any two of R², R³, R⁴, and R⁵ may collectively form a bridging group, provided that when the complex is of formula V or VIII, the bridging group is not a substituted sulfur atom or a substituted phosphorous atom;

10 Q represents a hydrocarbyl, chloride, iodide or bromide;

W represents a hydrocarbyl, chloride, iodide or bromide;

M represents Ni(II);

N represents nitrogen; O represents oxygen; S represents sulfur;

- 137 -

and Y is selected from the group consisting of a neutral Lewis acid capable of abstracting Q⁻ or W⁻ to form a weakly coordinating anion, a cationic Lewis acid whose counterion is a weakly coordinating anion, or a Bronsted acid whose conjugate base is a weakly coordinating anion.

5

146. The process of Claim 145, wherein the complex is of the formula II.

10 147. The process of Claim 145, wherein the complex is of the formula V.

148. The process of Claim 145, wherein the complex is of the formula VIII.

15

149. The process of Claim 145, wherein the complex is of the formula XV.

150. The process of Claim 145, wherein the complex is of the formula XVI.

20

151. The process of Claim 145 wherein R¹ and R⁶ are phenyl.

152. The process of Claim 145, wherein the process is conducted in an inert solvent.

25

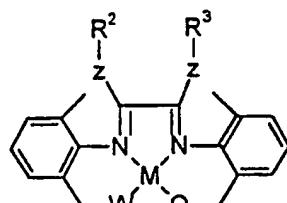
153. The process of Claim 152, wherein the inert solvent is selected from linear or branched hydrocarbons, toluene, or mixtures thereof.

- 138 -

154. The process of Claim 145 wherein the temperature is between about 25 °C and about 150 °C, and the pressure is between about 1 atm and about 1000 atm.

5 155. A process for the copolymerization of ethylene and a comonomer of the formula $\text{CH}_2=\text{CH}(\text{CH}_2)_n\text{CO}_2\text{R}^1$ which comprises contacting ethylene and a comonomer of the formula $\text{CH}_2=\text{CH}(\text{CH}_2)_n\text{CO}_2\text{R}^1$ with the complex formed by combining a first compound Y, which is selected from a neutral Lewis acid capable of abstracting Q⁻ or W⁻ to form a weakly coordinating anion, a cationic Lewis acid whose counterion is a weakly coordinating anion, or a Bronsted acid whose conjugate base is a weakly coordinating anion; with a second compound of the formula XXIV:

10



XXIV

15 wherein R¹ is hydrogen, hydrocarbyl, substituted hydrocarbyl, fluoroalkyl or silyl;

n is an integer greater than 3;

R² and R³ are each independently hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl, or may collectively form a bridging hydrocarbyl, bridging substituted hydrocarbyl, or a substituted silicon atom;

Q is alkyl, chloride, iodide or bromide;

W is alkyl, chloride, iodide or bromide;

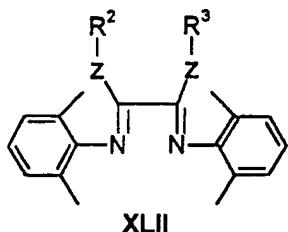
N is nitrogen;

Z is sulfur or oxygen; and

25 M is Ni(II).

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156. A process for the polymerization of ethylene and a comonomer of the formula $\text{CH}_2=\text{CH}(\text{CH}_2)_n\text{CO}_2\text{R}^1$, which comprises contacting ethylene and a comonomer of the formula $\text{CH}_2=\text{CH}(\text{CH}_2)_n\text{CO}_2\text{R}^1$ with the complex formed by the reaction product of (a) a compound of formula XLII ; (b) a suitable nickel(0) precursor, and (c) a Bronsted acid whose conjugate base is a weakly coordinating anion,



10 wherein R^1 is hydrogen, hydrocarbyl, substituted hydrocarbyl, fluoroalkyl or silyl;

 n is an integer greater than 3;

R^2 and R^3 are each independently hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl, or may collectively form a bridging hydrocarbyl, bridging substituted hydrocarbyl, or a substituted silicon atom;

15 N is nitrogen; and

 Z is sulfur or oxygen.

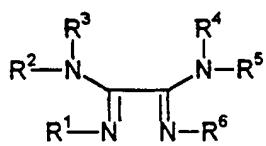
157. A process for the polymerization of olefins comprising

20 contacting one or more monomers of the formula $\text{RCH}=\text{CHR}^8$ with a binucleating or multinucleating ligand complexed to a Group 8-10 transition metal M and one or more Lewis acids, wherein the Lewis acid or acids are bound to one or more heteroatoms which are π -conjugated to the donor atom or atoms bound to the transition metal M; and R and R^8 each,

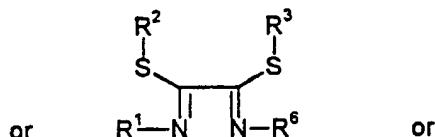
25 independently, represent a hydrogen, a hydrocarbyl, or a fluoroalkyl, and may be linked to form a cyclic olefin.

- 140 -

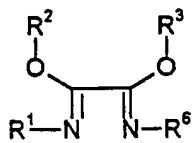
158. The process of Claim 157 wherein the binucleating or multinucleating ligand is selected from:



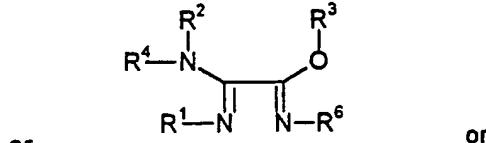
III



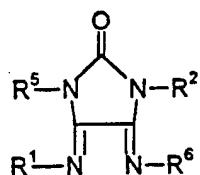
VI



IX



XVII



XVIII

5

and M is Ni(II), Pd(II), Co(II) or Fe(II), wherein R¹ and R⁶ each, independently, represent hydrocarbyl, substituted hydrocarbyl, or silyl; R², R³, R⁴ and R⁵ each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl; in addition, any two of R², R³, R⁴, and R⁵ may collectively form a bridging group, provided that when the ligand is of formula VI or IX, the bridging group is not a substituted sulfur atom or a substituted phosphorous atom; and

10

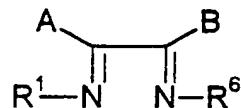
- 141 -

N represents nitrogen; O represents oxygen; and S represents sulfur.

159. The process of Claim 157 wherein the Lewis acid is a boron
5 or aluminum containing Lewis acid.

160. The process of Claim 158 wherein the Lewis acid is a boron
or aluminum containing Lewis acid.

10 161. A composition comprising (a) a Group 8-10 transition metal M,
(b) one or more Lewis acids, and (c) a binucleating or multinucleating
compound of the formula X :



X

15 wherein the Lewis acid or acids are bound to one or more
heteroatoms which are π -conjugated to the donor atom or atoms bound to
the transition metal M;

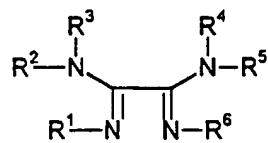
R¹ and R⁶ each, independently, represent sterically hindered aryl;

N represents nitrogen;

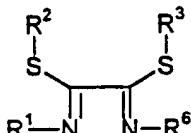
20 A and B are each, independently, a heteroatom connected mono-
radical wherein the connected heteroatom is selected from Group 15 or 16;
in addition, A and B may be linked by a bridging group.

25 162. The composition as in Claim 161 wherein the transition metal
is Ni(II), the Lewis acid is a boron or aluminum containing Lewis acid and
the binucleating or multinucleating compound of formula X is selected from

- 142 -



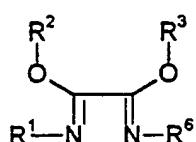
III



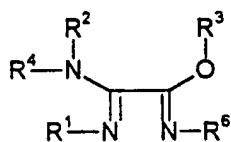
or

VI

or



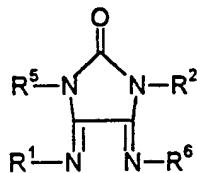
IX



XVII

or

or



XVIII

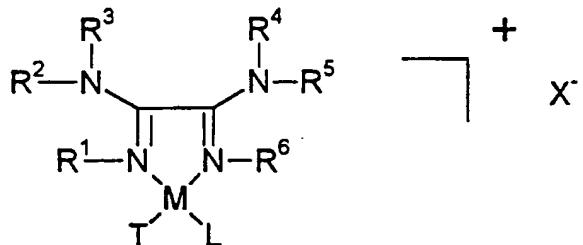
wherein R^1 and R^6 each, independently, represent sterically hindered aryl;

5 R^2 , R^3 , R^4 and R^5 each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl; in addition, any two of R^2 , R^3 , R^4 , and R^5 may collectively form a bridging group, provided that when the compound is of formula VI or IX, the bridging group is not a substituted sulfur atom or a substituted phosphorous atom;

10 N represents nitrogen; O represents oxygen; and S represents sulfur.

163. A compound of the formula I:

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I

wherein R¹ and R⁶ each, independently, represent hydrocarbyl, substituted hydrocarbyl, or silyl;

5 R², R³, R⁴ and R⁵ each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl; in addition, any two of R², R³, R⁴, and R⁵ may collectively form a bridging group;

T represents a hydrogen or hydrocarbyl;

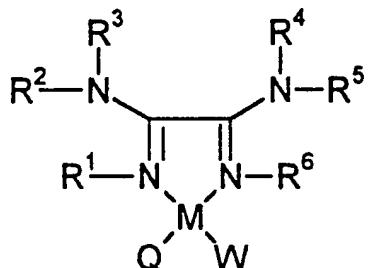
L represents a mono-olefin or a neutral Lewis base wherein the coordinated atom is nitrogen, oxygen, or sulfur;

10 M represents Ni(II), Pd(II), Co(II), or Fe(II);

N represents nitrogen; and

X⁻ is a weakly coordinating anion.

164. A compound of formula II:



II

15

wherein R¹ and R⁶ each, independently, represent a sterically hindered aryl;

- 144 -

R^2 , R^3 , R^4 and R^5 each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl; in addition, any two of R^2 , R^3 , R^4 , and R^5 may collectively form a bridging group;

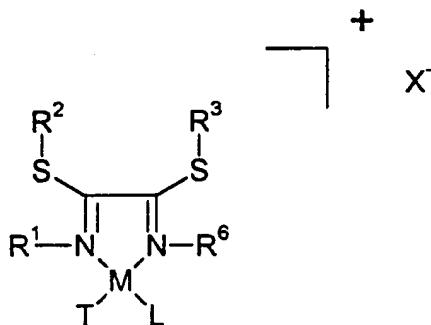
Q represents a hydrocarbyl, chloride, iodide or bromide;

5 W represents a hydrocarbyl, chloride, iodide or bromide;

N represents nitrogen; and

M represents $Ni(II)$, $Pd(II)$, $Co(II)$, or $Fe(II)$.

165. A compound of formula IV:



IV

10 wherein R^1 and R^6 each, independently, represent hydrocarbyl, substituted hydrocarbyl, or silyl;

R^2 and R^3 each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl, and, in addition, may collectively form a bridging hydrocarbyl, bridging substituted hydrocarbyl, or a substituted silicon atom;

15 T represents a hydrogen or hydrocarbyl;

L represents a mono-olefin or a neutral Lewis base wherein the coordinated atom is nitrogen, oxygen, or sulfur;

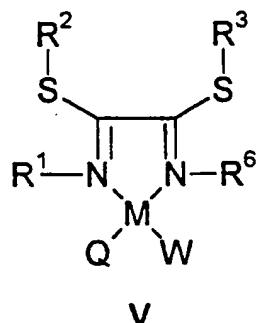
20 M represents $Ni(II)$, $Pd(II)$, $Co(II)$, or $Fe(II)$;

N represents nitrogen; and

X^- is a weakly coordinating anion.

- 145 -

166. A compound of formula V:



wherein R¹ and R⁶ each, independently, represent sterically hindered

5 aryl;

R² and R³ each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl, and, in addition, may collectively form a bridging hydrocarbyl, bridging substituted hydrocarbyl, or a substituted silicon atom;

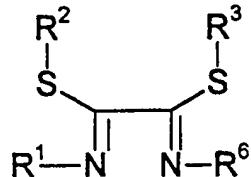
10 Q represents a hydrocarbyl, chloride, iodide or bromide;

W represents a hydrocarbyl, chloride, iodide or bromide;

N represents nitrogen; and

M represents Ni(II), Pd(II), Co(II), or Fe(II).

15 167. A compound of formula VI:



VI

wherein R¹ and R⁶ each, independently, represent a sterically

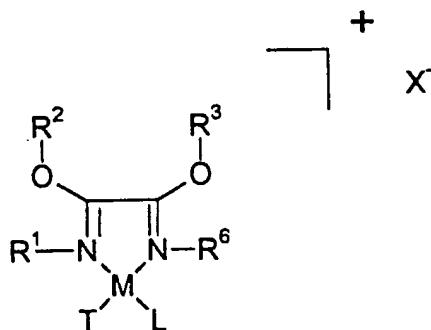
hindered aryl; and

- 146 -

R^2 and R^3 each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl, and, in addition, may collectively form a bridging hydrocarbyl, bridging substituted hydrocarbyl, or a substituted silicon atom.

5

168. A compound of the formula VII:



VII

wherein R^1 and R^6 each, independently, represent hydrocarbyl, substituted hydrocarbyl, or silyl;

10 R^2 and R^3 each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl, and, in addition, may collectively form a bridging hydrocarbyl, bridging substituted hydrocarbyl, or a substituted silicon atom;

T represents a hydrogen or hydrocarbyl;

15 L represents a mono-olefin or a neutral Lewis base wherein the coordinated atom is nitrogen, oxygen, or sulfur;

M represents Ni(II), Pd(II), Co(II), or Fe(II);

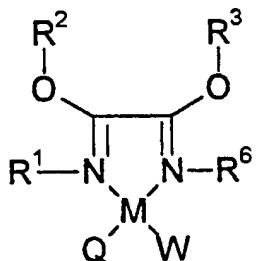
N represents nitrogen; and

X⁻ is a weakly coordinating anion.

20

169. A compound of formula VIII:

- 147 -



VIII

wherein R¹ and R⁶ each, independently, represent a sterically hindered aryl;

5 R² and R³ each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl, and, in addition, may collectively form a bridging hydrocarbyl, bridging substituted hydrocarbyl, or a substituted silicon atom;

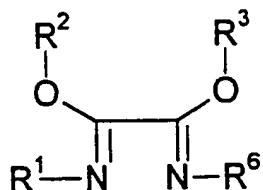
Q represents a hydrocarbyl, chloride, iodide or bromide;

W represents a hydrocarbyl, chloride, iodide or bromide;

10 N represents nitrogen; and

M represents Ni(II), Pd(II), Co(II), or Fe(II).

170. A compound of formula IX



IX

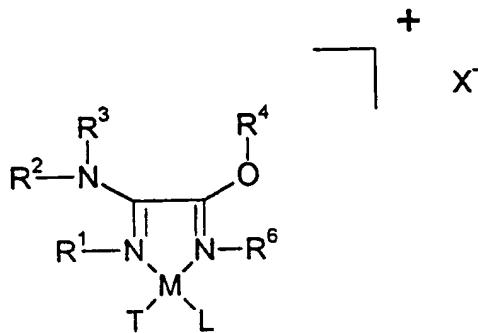
15 wherein R¹ and R⁶ each, independently, represent a sterically hindered aryl; and

R² and R³ each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl, and, in addition, may collectively form a

- 148 -

bridging hydrocarbyl, bridging substituted hydrocarbyl, or a substituted silicon atom.

171. A compound of the formula XIII:



XIII

5

wherein R¹ and R⁶ each, independently, represent hydrocarbyl, substituted hydrocarbyl, or silyl;

10 R², R³, and R⁴ each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl, and, in addition, any two of R², R³, and R⁴ may collectively form a bridging group;

T represents a hydrogen or hydrocarbyl;

15 L represents a mono-olefin or a neutral Lewis base wherein the coordinated atom is nitrogen, oxygen, or sulfur;

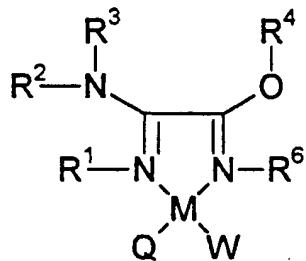
M represents Ni(II), Pd(II), Co(II), or Fe(II);

N represents nitrogen; and

X⁻ is a weakly coordinating anion.

172. A compound of formula XV:

- 149 -



XV

wherein R¹ and R⁶ each, independently, represent a sterically hindered aryl;

R², R³, and R⁴ each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl, and, in addition, any two of R², R³, and R⁴ may collectively form a bridging group;

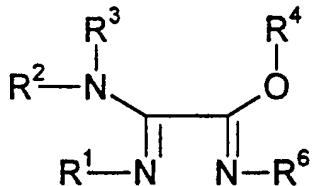
Q represents a hydrocarbyl, chloride, iodide or bromide;

W represents a hydrocarbyl, chloride, iodide or bromide;

N represents nitrogen; and

M represents Ni(II), Pd(II), Co(II), or Fe(II).

173. A compound of formula **XVII**



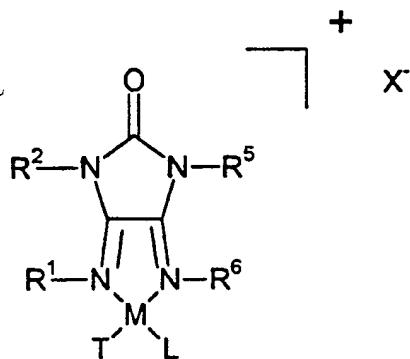
XVII

wherein R¹ and R⁶ each, independently, represent a sterically hindered aryl; and

R², R³, and R⁴ each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl, and, in addition, any two of R², R³, and R⁴ may collectively form a bridging group.

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174. A compound of the formula XIV:



XIV

wherein R¹ and R⁶ each, independently, represent hydrocarbyl, substituted hydrocarbyl, or silyl;

5 R² and R⁵ each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl;

T represents a hydrogen or hydrocarbyl;

L represents a mono-olefin or a neutral Lewis base wherein the coordinated atom is nitrogen, oxygen, or sulfur;

10 M represents Ni(II), Pd(II), Co(II), or Fe(II);

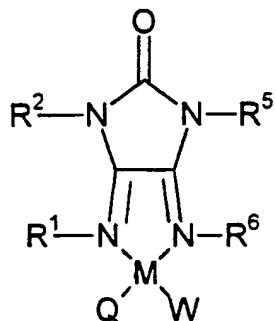
N represents nitrogen; and

X⁻ is a weakly coordinating anion.

175. A compound of formula XVI:

15

- 151 -



XVI

wherein R¹ and R⁶ each, independently, represent a sterically hindered aryl;

5 R² and R⁵ each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl;

Q represents a hydrocarbyl, chloride, iodide or bromide;

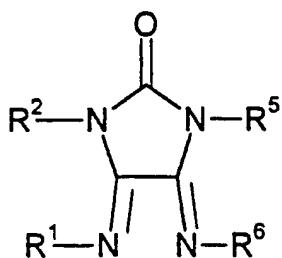
W represents a hydrocarbyl, chloride, iodide or bromide;

N represents nitrogen; and

M represents Ni(II), Pd(II), Co(II), or Fe(II).

10

176. A compound of formula XVIII:



XVIII

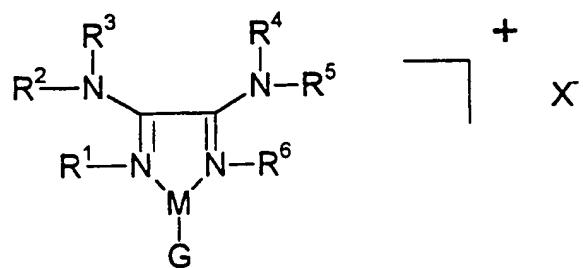
- 152 -

wherein R¹ and R⁶ each, independently, represent a sterically hindered aryl;

R² and R⁵ each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl.

5

177. A compound of the formula XIX:



wherein R¹ and R⁶ each, independently, represent hydrocarbyl, substituted hydrocarbyl, or silyl;

10 R², R³, R⁴ and R⁵ each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl; in addition, any two of R², R³, R⁴, and R⁵ may collectively form a bridging group;

G is a π -allyl or π -benzyl group;

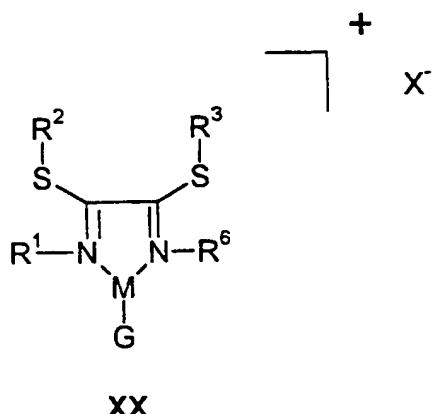
M represents Ni(II) or Pd(II);

15 N represents nitrogen; and

X⁻ is a weakly coordinating anion.

178. A compound of formula XX:

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wherein R¹ and R⁶ each, independently, represent hydrocarbyl, substituted hydrocarbyl, or silyl;

5 R² and R³ each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl, and, in addition, may collectively form a bridging hydrocarbyl, bridging substituted hydrocarbyl, or a substituted silicon atom;

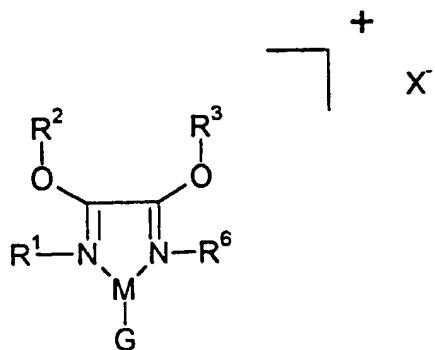
G is a π -allyl or π -benzyl;

M represents Ni(II), Pd(II);

10 N represents nitrogen; and

X⁻ is a weakly coordinating anion.

179. A compound of the formula XXI:



XXI

- 154 -

wherein R¹ and R⁶ each, independently, represent hydrocarbyl, substituted hydrocarbyl, or silyl;

R² and R³ each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl, and, in addition, may collectively form a bridging hydrocarbyl, bridging substituted hydrocarbyl, or a substituted silicon atom;

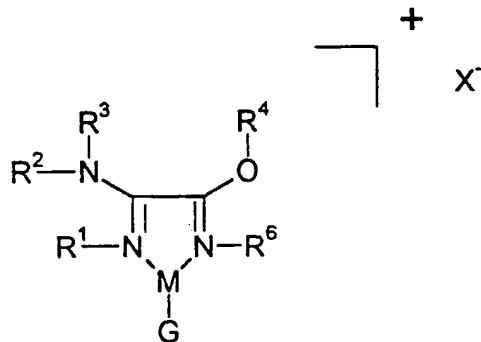
G is a π -allyl or π -benzyl;

M represents Ni(II), Pd(II);

N represents nitrogen; and

X⁻ is a weakly coordinating anion.

180. A compound of the formula XXII:



XXII

wherein R¹ and R⁶ each, independently, represent hydrocarbyl, substituted hydrocarbyl, or silyl;

R², R³, and R⁴ each, independently, represent a hydrogen, hydrocarbyl, substituted hydrocarbyl, or silyl, and, in addition, may collectively form a bridging group;

G is a π -allyl or π -benzyl;

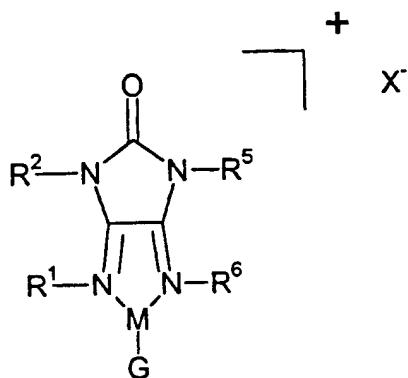
M represents Ni(II) or Pd(II);

N represents nitrogen; and

X⁻ is a weakly coordinating anion.

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181. A compound of the formula **XXIII**:



XXIII

wherein R¹ and R⁶ each, independently, represent hydrocarbyl,

5 substituted hydrocarbyl, or silyl;

R² and R⁵ each, independently, represent a hydrogen, hydrocarbyl,

substituted hydrocarbyl, or silyl;

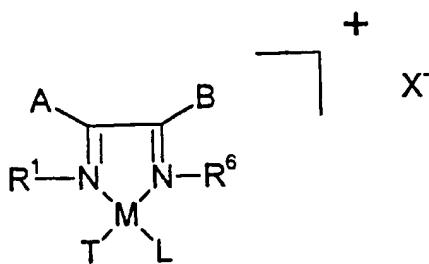
G is a π -allyl or π -benzyl;

M represents Ni(II), Pd(II);

10 N represents nitrogen; and

X⁻ is a weakly coordinating anion.

182. A compound of formula **XXIV**,



XXIV

15

- 156 -

wherein R¹ and R⁶ each, independently, represent hydrocarbyl, substituted hydrocarbyl, or silyl;

5 A and B are each, independently, a heteroatom connected mono-radical wherein the connected heteroatom is selected from Group 15 or 16; in addition, A and B may be linked by a bridging group;

T represents a hydrogen or a hydrocarbyl;

L represents a mono-olefin or a neutral Lewis base wherein the coordinated atom is nitrogen, oxygen, or sulfur;

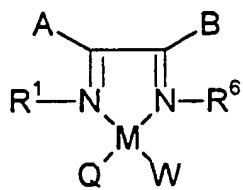
10 M represents Ni(II) or Pd(II);

N represents nitrogen; and

X⁻ is a weakly coordinating anion.

183. A compound of formula XXIX,

15



XXIX

wherein R¹ and R⁶ each, independently, represent a sterically hindered aryl;

20 A and B are each, independently, a heteroatom connected mono-radical wherein the connected heteroatom is selected from Group 15 or 16; in addition, A and B may be linked by a bridging group;

Q represents a hydrocarbyl, chloride, iodide or bromide;

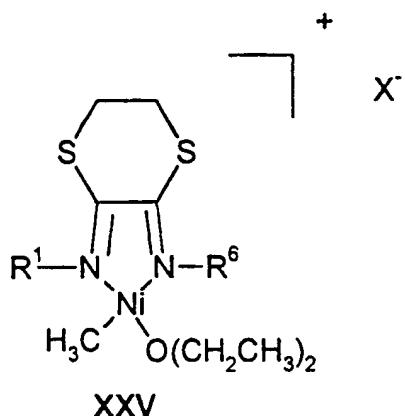
W represents a hydrocarbyl, chloride, iodide or bromide;

25 M represents Ni(II) or Pd(II); and

N represents nitrogen.

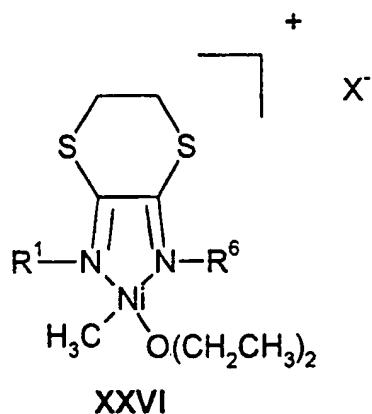
- 157 -

184. The compound of Claim 165 having the formula **XXV**:



wherein R¹ and R⁶ are 2,6-dimethylphenyl;
5 and X⁻ is a weakly coordinating anion.

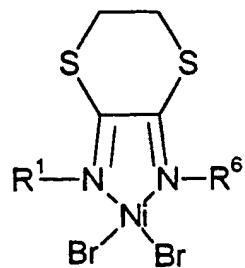
185. The compound of Claim 165 having the formula **XXVI**:



wherein R¹ and R⁶ are 2,6-diisopropylphenyl;
10 and X⁻ is a weakly coordinating anion.

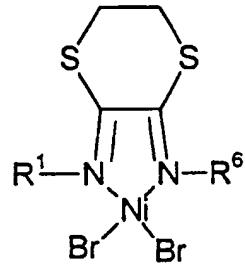
186. The compound of Claim 166 having the formula **XXVII**,

- 158 -



wherein R¹ and R⁶ are 2,6-dimethylphenyl.

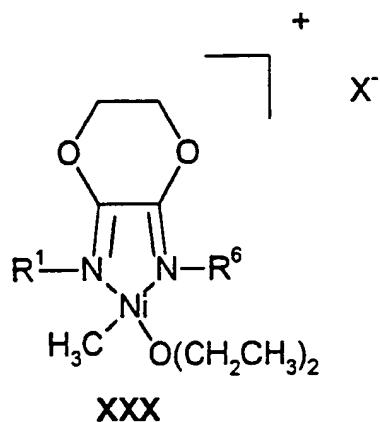
187. The compound of Claim 166 having the formula XXVIII,



5

wherein R¹ and R⁶ are 2,6-diisopropylphenyl.

188. The compound of Claim 168 having the formula XXX,

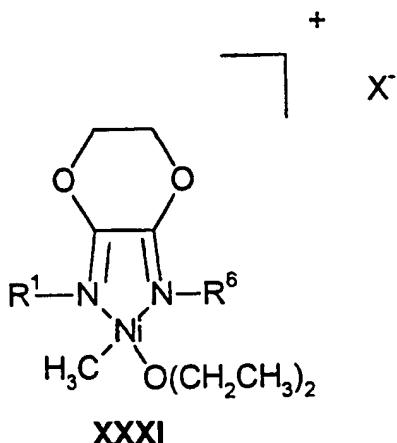


10 wherein R¹ and R⁶ are 2,6-dimethylphenyl;

- 159 -

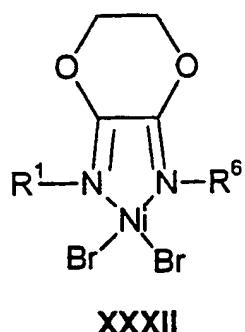
and X^- is a weakly coordinating anion.

189. The compound of Claim 168 having the formula XXXI,



5 wherein R^1 and R^6 are 2,6-diisopropylphenyl;
and X^- is a weakly coordinating anion.

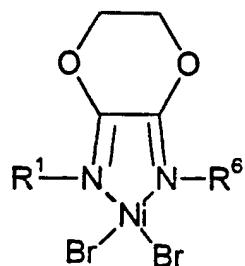
190. The compound of Claim 169 having the formula XXXII,



10 wherein R^1 and R^6 are 2,6-dimethylphenyl.

191. The compound of Claim 169 having the formula XXXIII,

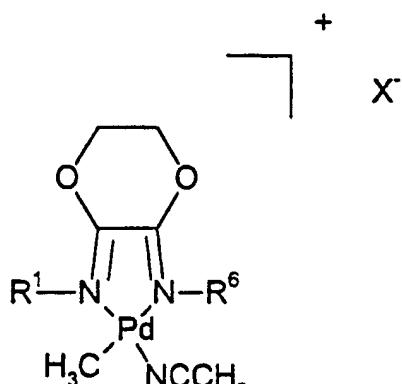
- 160 -



XXXIII

wherein R^1 and R^6 are 2,6-diisopropylphenyl.

192. The compound of Claim 168 having the formula XXXIV,



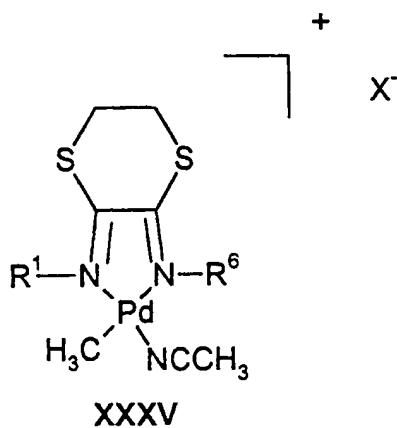
XXXIV

5

wherein R^1 and R^6 are 2,6-diisopropylphenyl;
and X^- is a weakly coordinating anion.

193. The compound of Claim 165 having the formula XXXV,

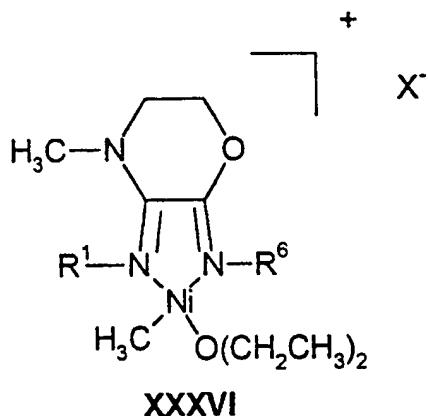
- 161 -



wherein R^1 and R^6 are 2,6-diisopropylphenyl;
and X^- is a weakly coordinating anion.

5

194. The compound of Claim 171 having the formula **XXXVI**,

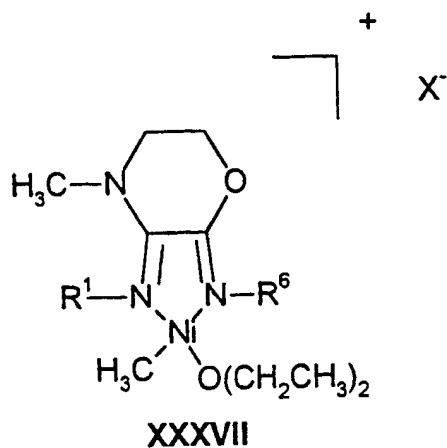


wherein R^1 and R^6 are 2,6-dimethylphenyl;
and X^- is a weakly coordinating anion.

10

195. The compound of Claim 171 having the formula **XXXVII**,

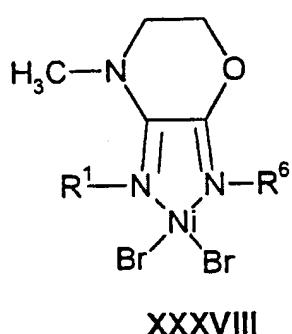
- 162 -



wherein R¹ and R⁶ are 2,6-diisopropylphenyl;
and X⁻ is a weakly coordinating anion.

5

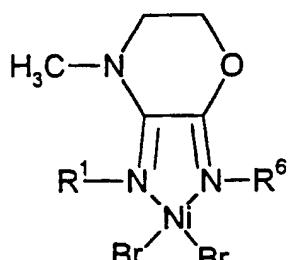
196. The compound of Claim 172 having the formula **XXXVIII**,



wherein R¹ and R⁶ are 2,6-dimethylphenyl.

197. The compound of Claim 172 having the formula **XXXIX**,

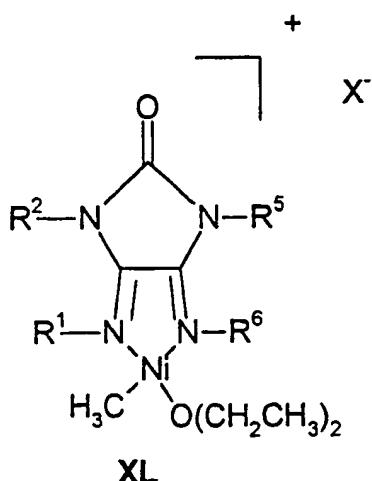
- 163 -



XXXIX

wherein R¹ and R⁶ are 2,6-diisopropylphenyl.

198. The compound of Claim 174 having the formula XL,

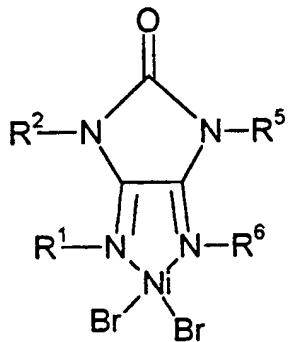


5

wherein R¹, R², R⁵ and R⁶ are 2,6-dimethyl-4-methoxyphenyl;
and X⁻ is a weakly coordinating anion.

199. The compound of Claim 175 having the formula XLI,

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XLI

wherein R¹, R², R⁵ and R⁶ are 2,6-dimethyl-4-methoxyphenyl.

200. A polymer composition comprised of monomer units derived
 5 from ethylene and from 0.1 to 50 wt % of a comonomer of the formula
 $\text{CH}_2=\text{CH}(\text{CH}_2)_n\text{CO}_2\text{R}^1$, wherein R¹ is hydrogen, hydrocarbyl, substituted
 hydrocarbyl, fluoroalkyl or silyl, and n is an integer between 3 and 18; in
 addition to the branches attributable to the incorporation of said
 comonomer, alkyl branches are present in said polymer composition,
 10 wherein between 5 and 50 alkyl branches per 1000 carbon atoms are
 present and the majority of alkyl branches are methyl branches.

201. A composition comprising an ester containing semicrystalline
 15 copolyethylene with 5 to 30 alkyl branches/1000 carbon atoms, wherein the
 majority of alkyl branches are methyl branches, and an average of from
 about 1 to 50 ester terminated branches per chain resulting from ester
 comonomer incorporation.

202. A composition comprising an olefin containing semicrystalline
 20 copolyethylene with 5 to 30 alkyl branches/1000 carbon atoms, wherein the
 majority of alkyl branches are methyl branches, and an average of from
 about 1 to 50 olefin terminated branches per chain resulting from linear
 diene comonomer incorporation.

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203. The compound of Claim 164 wherein M is Ni(II).

5 204. A composition comprising an ethylene homopolymer with
greater than 125 alkyl branches per 1000 carbon atoms.